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2019

Cracking the
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PREMIUM

5 full-length practice tests (4 in the book & 1 online)
with complete answer explanations

- Comprehensive review of key AP Chemistry concepts
- Targeted strategies for every section of the exam
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EXAM
PREMIUM**

2019 Edition

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- If you're still choosing between colleges, use our searchable rankings of *The Best 384 Colleges* to find out more information about your dream school.
- Access comprehensive study guides and a variety of printable resources, including: answer bubble sheets, the periodic table, and a list of equations to review
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Look For These Icons Throughout The Book



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Part I

Using This Book to Improve Your AP Score

PREVIEW ACTIVITY: YOUR KNOWLEDGE, YOUR EXPECTATIONS

Your route to a high score on the AP Chemistry Exam depends a lot on how you plan to use this book. Please respond to the following questions.

1. Rate your level of confidence about your knowledge of the content tested by the AP Chemistry Exam:
 - A. Very confident—I know it all
 - B. I'm pretty confident, but there are topics for which I could use help
 - C. Not confident—I need quite a bit of support
 - D. I'm not sure
2. Circle your goal score for the AP Chemistry Exam.

5	4	3	2	1	I'm not sure yet
---	---	---	---	---	------------------
3. What do you expect to learn from this book? Circle all that apply to you.
 - A. A general overview of the test and what to expect
 - B. Strategies for how to approach the test
 - C. The content tested by this exam
 - D. I'm not sure yet

YOUR GUIDE TO USING THIS BOOK

This book is organized to provide as much—or as little—support as you need, so you can use this book in whatever way will be most helpful to improving your score on the AP Chemistry Exam.

- The remainder of **Part I** will provide guidance on how to use this book and help you determine your strengths and weaknesses
- **Part II** of this book contains Practice Test 1 and its answers and explanations. (Bubble sheets can be found in the very back of the book for easy tear-out.) We strongly recommend that you take this test before going any further, in order to realistically determine:
 - your starting point right now
 - which question types you're ready for and which you might need to practice
 - which content topics you are familiar with and which you will want to carefully review

Once you have nailed down your strengths and weaknesses with regard to this exam, you can focus your test preparation, build a study plan, and be efficient with your time.

- **Part III** of this book will
 - provide information about the structure, scoring, and content of the AP Chemistry Exam
 - help you to make a study plan
 - point you toward additional resources
- **Part IV** of this book will explore
 - how to attack multiple-choice questions
 - how to write high-scoring free-response answers
 - how to manage your time to maximize the number of points available to you
- **Part V** of this book covers the content you need for your exam.

- **Parts VI and VII** of this book contain Practice Tests 2, 3, and 4, their answers and explanations, and a scoring guide. (Bubble sheets can be found in the very back of the book for easy tear-out.) If you skipped Practice Test 1, we recommend that you do both (with at least a day or two between them) so that you can compare your progress between the two. Additionally, this will help to identify any external issues: If you get a certain type of question wrong both times, you probably need to review it. If you only got it wrong once, you may have run out of time or been distracted by something. In either case, this will allow you to focus on the factors that caused the discrepancy in scores and to be as prepared as possible on the day of the test.

You may choose to use some parts of this book over others, or you may work through the entire book. This will depend on your needs and how much time you have. Let's now look how to make this determination.

HOW TO BEGIN

1. Take a Test

Before you can decide how to use this book, you need to take a practice test. Doing so will give you insight into your strengths and weaknesses, and the test will also help you make an effective study plan. If you're feeling test-phobic, remind yourself that a practice test is a tool for diagnosing yourself—it's not how well you do that matters but how you use information gleaned from your performance to guide your preparation.

So, before you read further, take AP Chemistry Practice Test 1 starting at [this page](#) of this book. Be sure to do so in one sitting, following the instructions that appear before the test.

2. Check Your Answers

Using the answer key on [this page](#), count how many multiple-choice questions you got right and how many you missed. Don't worry about the explanations for now, and don't worry about why you missed questions. We'll get to that soon.

3. Reflect on the Test

After you take your first test, respond to the following questions:

- How much time did you spend on the multiple-choice questions?
- How much time did you spend on each long form free-response question? What about each short form free-response question?
- How many multiple-choice questions did you miss?
- Do you feel you had the knowledge to address the subject matter of the free-response questions?
- Do you feel you wrote well organized, thoughtful answers to the free-response questions?
- Circle the content areas that were most challenging for you and draw a line through the ones in which you felt confident/did well.
 - **Big Idea #1:** Atoms, Elements, and the Building Blocks of Matter
 - **Big Idea #2:** Bonding and Phases
 - **Big Idea #3:** Chemical Reactions, Energy Changes, and Redox Reactions
 - **Big Idea #4:** Chemical Reactions and their Rates
 - **Big Idea #5:** Laws of Thermodynamics and Changes in Matter

- **Big Idea #6:** Equilibrium, Acids and Bases, Titrations and Solubility

4. Read Part III of this Book and Complete the Self-Evaluation

As discussed previously, Part III will provide information on how the test is structured and scored. As you read Part III, re-evaluate your answers to the questions on the previous page. You will then be able to make a study plan, based on your needs and time available, that will allow you to use this book most effectively.

5. Engage with Parts IV and V as Needed

Notice the word *engage*. You'll get more out of this book if you use it intentionally than if you read it passively, hoping for an improved score through osmosis.

The strategy chapters will help you think about your approach to the question types on this exam. Part IV will open with a reminder to think about how you approach questions now and then close with a reflection section asking you to think about how/whether you will change your approach in the future.

The content chapters are designed to provide a review of the content tested on the AP Chemistry Exam, including the level of detail you need to know and how the content is tested. You will have the opportunity to assess your mastery of the content of each chapter through test-appropriate questions.

6. Take Test 2 and Assess Your Performance

Once you feel you have developed the strategies you need and gained the knowledge you lacked, you should take Test 2, which starts at [this page](#) of this book. You should do so in one sitting, following the instructions at the beginning of the test.

When you are done, check your answers to the multiple-choice sections on [this page](#). See if a teacher will read your answers to the free-response questions and provide feedback.

Once you have taken the test, reflect on what areas you still need to work on, and revisit the chapters in this book that address those deficiencies. Through this type of reflection and engagement, you will continue to improve.

7. Keep Working

After you have revisited certain chapters in this book, continue the process of testing, reflecting, and engaging with the second test. Each time, consider what additional work you need to do and how you will change your strategic approach to different parts of the test.

Part II

Practice Test 1

[Click here](#) to download and print the PDF version of this exercise.

AP[®] Chemistry Exam

SECTION I: Multiple-Choice Questions

DO NOT OPEN THIS BOOKLET UNTIL YOU ARE TOLD TO DO SO.

At a Glance

Total Time

1 hour and 30 minutes

Number of Questions

60

Percent of Total Grade

50%

Writing Instrument

Pencil required

Instructions

Section I of this examination contains 60 multiple-choice questions. Fill in only the ovals for numbers 1 through 60 on your answer sheet.

CALCULATORS MAY NOT BE USED IN THIS PART OF THE EXAMINATION.

Indicate all of your answers to the multiple-choice questions on the answer sheet. No credit will be given for anything written in this exam booklet, but you may use the booklet for notes or scratch work. After you have decided which of the suggested answers is best, completely fill in the corresponding oval on the answer sheet. Give only one answer to each question. If you change an answer, be sure that the previous mark is erased completely. Here is a sample question and answer.

Sample Question

Chicago is a

- (A) state
- (B) city
- (C) country
- (D) continent

Sample Answer

☐ (A) ☒ (B) ☐ (C) ☐ (D)

Use your time effectively, working as quickly as you can without losing accuracy. Do not spend too much time on any one question. Go on to other questions and come back to the ones you have not answered if you have time. It is not expected that everyone will know the answers to all the multiple-choice questions.

About Guessing

Many candidates wonder whether or not to guess the answers to questions about which they are not certain. Multiple-choice scores are based on the number of questions answered correctly. Points are not deducted for incorrect answers, and no points are awarded for unanswered questions. Because points are not deducted for incorrect answers, you are encouraged to answer all multiple-choice questions. On any questions you do not know the answer to, you should eliminate as many choices as you can, and then select the best answer among the remaining choices.

Section I

CHEMISTRY SECTION I

Time—1 hour and 30 minutes

INFORMATION IN THE TABLE BELOW AND ON THE FOLLOWING PAGES MAY BE USEFUL IN ANSWERING THE QUESTIONS IN THIS SECTION OF THE EXAMINATION.

DO NOT DETACH FROM BOOK.

PERIODIC TABLE OF THE ELEMENTS

PERIODIC TABLE OF THE ELEMENTS																	2
1 H 1.008																	2 He 4.00
3 Li 6.94	4 Be 9.01											5 B 10.81	6 C 12.01	7 N 14.01	8 O 16.00	9 F 19.00	10 Ne 20.18
11 Na 22.99	12 Mg 24.30											13 Al 26.98	14 Si 28.09	15 P 30.97	16 S 32.06	17 Cl 35.45	18 Ar 39.95
19 K 39.10	20 Ca 40.08	21 Sc 44.96	22 Ti 47.90	23 V 50.94	24 Cr 52.00	25 Mn 54.94	26 Fe 55.85	27 Co 58.93	28 Ni 58.69	29 Cu 63.55	30 Zn 65.39	31 Ga 69.72	32 Ge 72.59	33 As 74.92	34 Se 78.96	35 Br 79.90	36 Kr 83.80
37 Rb 85.47	38 Sr 87.62	39 Y 88.91	40 Zr 91.22	41 Nb 92.91	42 Mo 95.94	43 Tc (98)	44 Ru 101.1	45 Rh 102.91	46 Pd 106.42	47 Ag 107.87	48 Cd 112.41	49 In 114.82	50 Sn 118.71	51 Sb 121.75	52 Te 127.60	53 I 126.91	54 Xe 131.29
55 Cs 132.91	56 Ba 137.33	57 *La 138.91	72 Hf 178.49	73 Ta 180.95	74 W 183.85	75 Re 186.21	76 Os 190.2	77 Ir 192.2	78 Pt 195.08	79 Au 196.97	80 Hg 200.59	81 Tl 204.38	82 Pb 207.2	83 Bi 208.98	84 Po (209)	85 At (210)	86 Rn (222)
87 Fr (223)	88 Ra 226.02	89 †Ac 227.03	104 Rf (261)	105 Db (262)	106 Sg (266)	107 Bh (264)	108 Hs (277)	109 Mt (268)	110 Ds (271)	111 Rg (272)							

*Lanthanide Series

58 Ce 140.12	59 Pr 140.91	60 Nd 144.24	61 Pm (145)	62 Sm 150.4	63 Eu 151.97	64 Gd 157.25	65 Tb 158.93	66 Dy 162.50	67 Ho 164.93	68 Er 167.26	69 Tm 168.93	70 Yb 173.04	71 Lu 174.97
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†Actinide Series

90 Th 232.04	91 Pa 231.04	92 U 238.03	93 Np (237)	94 Pu (244)	95 Am (243)	96 Cm (247)	97 Bk (247)	98 Cf (251)	99 Es (252)	100 Fm (257)	101 Md (258)	102 No (259)	103 Lr (262)
---------------------------	---------------------------	--------------------------	--------------------------	--------------------------	--------------------------	--------------------------	--------------------------	--------------------------	--------------------------	---------------------------	---------------------------	---------------------------	---------------------------

ADVANCED PLACEMENT CHEMISTRY EQUATIONS AND CONSTANTS

Throughout the test the following symbols have the definitions specified unless otherwise noted.

L, mL = liter(s), milliliter(s)
 g = gram(s)
 nm = nanometer(s)
 atm = atmosphere(s)

mm Hg = millimeters of mercury
 J, kJ = joule(s), kilojoule(s)
 V = volt(s)
 mol = mole(s)

ATOMIC STRUCTURE

$$E = h\nu$$

$$c = \lambda\nu$$

E = energy
 ν = frequency
 λ = wavelength

Planck's constant, $h = 6.626 \times 10^{-34} \text{ J s}$

Speed of light, $c = 2.998 \times 10^8 \text{ m s}^{-1}$

Avogadro's number = $6.022 \times 10^{23} \text{ mol}^{-1}$

Electron charge, $e = -1.602 \times 10^{-19} \text{ coulomb}$

EQUILIBRIUM

$$K_c = \frac{[C]^c[D]^d}{[A]^a[B]^b}, \text{ where } aA + bB \rightleftharpoons cC + dD$$

$$K_p = \frac{(P_C)^c(P_D)^d}{(P_A)^a(P_B)^b}$$

$$K_a = \frac{[H^+][A^-]}{[HA]}$$

$$K_b = \frac{[OH^-][HB^+]}{[B]}$$

$$K_w = [H^+][OH^-] = 1.0 \times 10^{-14} \text{ at } 25^\circ\text{C}$$

$$= K_a \times K_b$$

$$\text{pH} = -\log[H^+], \text{ pOH} = -\log[OH^-]$$

$$14 = \text{pH} + \text{pOH}$$

$$\text{pH} = \text{p}K_a + \log \frac{[A^-]}{[HA]}$$

$$\text{p}K_a = -\log K_a, \text{ p}K_b = -\log K_b$$

Equilibrium Constants

K_c (molar concentrations)

K_p (gas pressures)

K_a (weak acid)

K_b (weak base)

K_w (water)

KINETICS

$$\ln[A]_t - \ln[A]_0 = -kt$$

$$\frac{1}{[A]_t} - \frac{1}{[A]_0} = kt$$

$$t_{1/2} = \frac{0.693}{k}$$

k = rate constant

t = time

$t_{1/2}$ = half-life

GASES, LIQUIDS, AND SOLUTIONS

$$PV = nRT$$

$$P_A = P_{\text{total}} \times X_A, \text{ where } X_A = \frac{\text{moles A}}{\text{total moles}}$$

$$P_{\text{total}} = P_A + P_B + P_C + \dots$$

$$n = \frac{m}{M}$$

$$K = ^\circ\text{C} + 273$$

$$D = \frac{m}{V}$$

$$KE \text{ per molecule} = \frac{1}{2}mv^2$$

Molarity, M = moles of solute per liter of solution

$$A = abc$$

P = pressure

V = volume

T = temperature

n = number of moles

m = mass

M = molar mass

D = density

KE = kinetic energy

v = velocity

A = absorbance

a = molar absorptivity

b = path length

c = concentration

Gas constant, $R = 8.314 \text{ J mol}^{-1} \text{ K}^{-1}$

$$= 0.08206 \text{ L atm mol}^{-1} \text{ K}^{-1}$$

$$= 62.36 \text{ L torr mol}^{-1} \text{ K}^{-1}$$

$$1 \text{ atm} = 760 \text{ mm Hg}$$

$$= 760 \text{ torr}$$

STP = 273.15 K and 1.0 atm

Ideal gas at STP = 22.4 L mol⁻¹

THERMOCHEMISTRY/ ELECTROCHEMISTRY

$$q = mc\Delta T$$

$$\Delta S^\circ = \sum S^\circ \text{ products} - \sum S^\circ \text{ reactants}$$

$$\Delta H^\circ = \sum \Delta H_f^\circ \text{ products} - \sum \Delta H_f^\circ \text{ reactants}$$

$$\Delta G^\circ = \sum \Delta G_f^\circ \text{ products} - \sum \Delta G_f^\circ \text{ reactants}$$

$$\Delta G^\circ = \Delta H^\circ - T\Delta S^\circ$$

$$= -RT \ln K$$

$$= -nFE^\circ$$

$$I = \frac{q}{t}$$

q = heat

m = mass

c = specific heat capacity

T = temperature

S° = standard entropy

H° = standard enthalpy

G° = standard free energy

n = number of moles

E° = standard reduction potential

I = current (amperes)

q = charge (coulombs)

t = time (seconds)

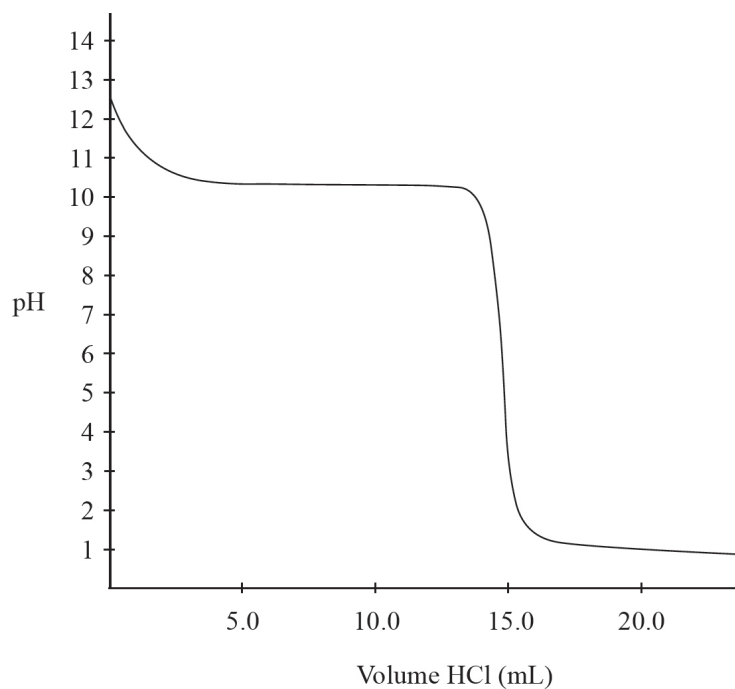
Faraday's constant, $F = 96,485$ coulombs per mole of electrons

$$1 \text{ volt} = \frac{1 \text{ joule}}{1 \text{ coulomb}}$$

1. In a saturated solution of Na_3PO_4 , $[\text{Na}^+] = 0.30\text{ M}$. What is the molar solubility of Na_3PO_4 ?
- (A) 0.10 M
(B) 0.30 M
(C) 0.60 M
(D) 0.90 M
2. When some LiCl is dissolved in water, the temperature of the water increases. This means that:
- (A) the strength of the intermolecular forces between the water molecules is stronger than the bond energy within the LiCl lattice
(B) the attraction of the lithium ions to the negative dipoles of the water molecules is weaker than the attraction of the chloride ions to the positive dipoles of the water molecules
(C) breaking the bonds between the lithium and chloride ions is an exothermic process
(D) the strength of the ion-dipole attractions that are formed exceeds the lattice energy in LiCl

Use the following information to answer questions 3-6

A student titrates some 1.0 M HCl into 20.0 mL of methylamine (CH_3NH_2), a weak base which only accepts a single proton. The following titration curve results:



3. What is the concentration of the methylamine?

- (A) 0.50 *M*
- (B) 0.75 *M*
- (C) 1.0 *M*
- (D) 1.25 *M*

4. What is the approximate pK_b for methylamine?

- (A) 3.5
- (B) 5.5
- (C) 10.5
- (D) 12.5

5. The buffer region of this titration is located:

- (A) Below 3.0 mL
- (B) Between 3.0 mL and 14.0 mL

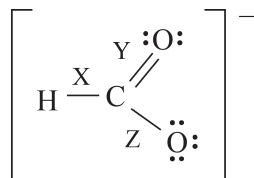
(C) Between 14.0 mL and 16.0 mL

(D) Above 16.0 mL

6. The methylamine is replaced by 20.0 mL of sodium hydroxide of an identical concentration. If the sodium hydroxide is titrated with the 1.0 *M* HCl, which of the following options accurately describes the pH levels at various points during the titration when compared to the pH levels at the same point in the HCl/methylamine titration?

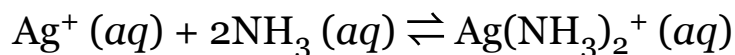
	Initial pH	Equivalence pH	Ending pH
(A)	lower	same	higher
(B)	higher	higher	same
(C)	same	higher	same
(D)	higher	lower	lower

7. The formate ion, HCO_2^- , is best represented by the Lewis diagram below. Each bond is labeled with a different letter.



What is the bond order for each bond?

	X	Y	Z
(A)	1	1	2
(B)	2	2	1
(C)	1	1.5	1.5
(D)	1.33	1.33	1.33

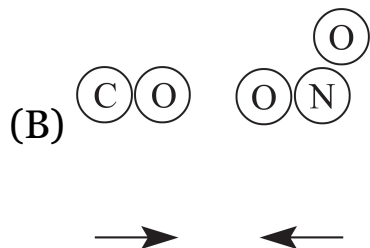
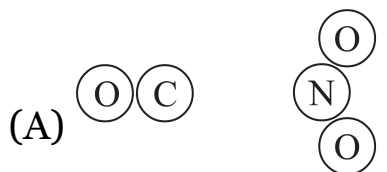


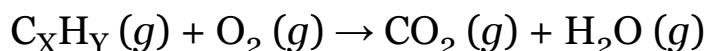
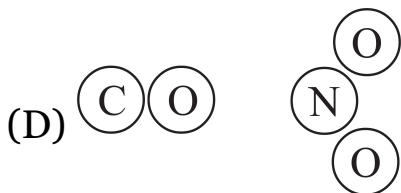
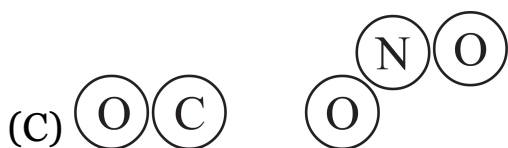
8. The reaction above is at equilibrium in a closed system. Which of the following will happen immediately when water is added?

- (A) The rate of the reverse reaction will increase.
- (B) Both ions will increase in concentration, while the NH_3 decreases in concentration.
- (C) The reaction will shift to the right.
- (D) Nothing will happen; adding water does not cause any changes to the equilibrium system.



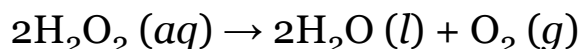
9. The two products above are created from a reaction involving CO and NO_2 . Which of the following options indicates a viable orientation of both reactant molecules prior to the collision? (Note: the arrows indicate the direction each molecule is moving prior to the collision.)





10. When the above unbalanced reaction occurs at STP, 1.5 L of CO_2 and 1.0 L of H_2O are created. What is the empirical formula of the hydrocarbon?

- (A) CH_2
 (B) C_2H_3
 (C) C_2H_5
 (D) C_3H_4



11. For the decomposition of hydrogen peroxide, which element (if any) is being reduced, and which is being oxidized?

	Oxidized	Reduced
(A)	Hydrogen	Oxygen
(B)	Oxygen	None
(C)	None	Hydrogen
(D)	Oxygen	Oxygen

12. Identical amounts of the four gases listed below are present in four separate balloons. At STP, which balloon size experiences the greatest deviation from the volume calculated using the Ideal Gas Law?

(A) H₂

(B) O₂

(C) N₂

(D) F₂

13. Which of the following reasons correctly explains one reason that increasing the temperature of a reaction increases its speed?

(A) All reactant molecules will have more kinetic energy.

(B) A larger percentage of reactant molecules will exceed the activation energy barrier.

(C) A higher percentage of molecular collisions will have the correct orientation to cause a reaction.

(D) The order of each reactant will increase.

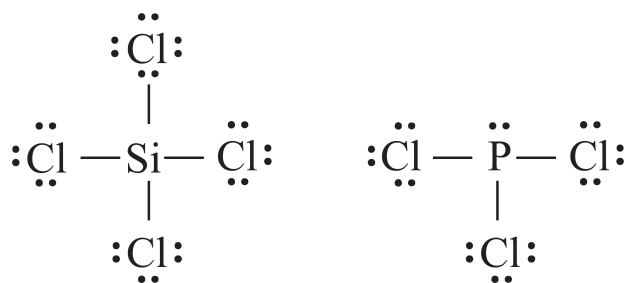
Use the following information to answer questions 14-16

The radius of atoms and ions is typically measured in Angstroms (Å), which is equivalent to 1×10^{-10} m. Below is a table of information for three different elements.

Element	Atomic Radius (Å)	Ionic Radius (Å)
Ne	0.38	N/A
P	0.98	1.00

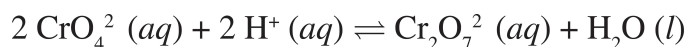
Zn	1.42	1.35
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14. The phosphorus ion is larger than a neutral phosphorus atom, yet a zinc ion is smaller than a neutral zinc atom. Which of the following statements best explains why?
- (A) The zinc atom has more protons than the phosphorus atom.
 - (B) The phosphorus atom has fewer valence electrons than the zinc atom.
 - (C) Phosphorus gains electrons when forming an ion, but zinc loses them.
 - (D) The valence electrons in zinc are further from the nucleus than those in phosphorus.
15. Neon has a smaller atomic radius than phosphorus because:
- (A) Unlike neon, phosphorus has electrons present in its third energy level.
 - (B) Phosphorus has more protons than neon, which increases the repulsive forces in the atom.
 - (C) The electrons in a neon atom are all found in a single energy level.
 - (D) Phosphorus can form anions, while neon is unable to form any ions.
16. Which of the following represents the correct electron configuration for the zinc ion, Zn^{2+} ?
- (A) $[\text{Ar}]3\text{d}^{10}$
 - (B) $[\text{Ar}]4\text{s}^23\text{d}^8$
 - (C) $[\text{Ar}]4\text{s}^24\text{d}^8$
 - (D) $[\text{Kr}]4\text{s}^23\text{d}^8$



17. The Lewis diagrams for SiCl_4 and PCl_3 are drawn above. What are the approximate bond angles between the terminal chlorine atoms in each structure?

	SiCl_4	PCl_3
(A)	90	90
(B)	109.5°	$< 109.5^\circ$
(C)	90°	109.5°
(D)	$< 109.5^\circ$	$> 90^\circ$



18. The above reaction is present at equilibrium in a beaker. A student stirs the mixture. What effect will this have on the reaction rates?
- (A) It will increase both the forward and reverse reaction rates.
- (B) It will increase the forward rate, but decrease the reverse rate.
- (C) It will have no effect on the forward rate, but it will decrease the reverse rate.
- (D) It will have no effect on either rate.
19. A sample of water originally at 25°C is heated to 75°C . As the temperature increases, the vapor pressure of the water is also observed to increase. Why?

- (A) Water molecules are more likely to have enough energy to break free of the intermolecular forces holding them together.
- (B) The covalent bonds between the hydrogen and oxygen atoms within individual water molecules are more likely to be broken.
- (C) The strength of the hydrogen bonding between different water molecules will increase until it exceeds the covalent bond energy within individual water molecules.
- (D) The electron clouds surrounding each water molecule are becoming less polarizable, weakening the intermolecular forces between them.
20. The enthalpy change for which of the following reactions would be equal to the enthalpy of formation for ethanol ($\text{CH}_3\text{CH}_2\text{OH}$)?
- (A) $\text{CH}_3 + \text{CH}_2 + \text{OH} \rightarrow \text{CH}_3\text{CH}_2\text{OH}$
- (B) $2\text{C} + 5\text{H} + \text{O} \rightarrow \text{CH}_3\text{CH}_2\text{OH}$
- (C) $4\text{C} + 6\text{H}_2 + \text{O}_2 \rightarrow 2\text{CH}_3\text{CH}_2\text{OH}$
- (D) $2\text{C} + 3\text{H}_2 + \frac{1}{2}\text{O}_2 \rightarrow \text{CH}_3\text{CH}_2\text{OH}$
21. A chemist wants to plate out 1.00 g of solid iron from a solution containing aqueous Fe^{2+} ions. Which of the following expressions will equal the amount of time, in seconds, it takes if a current of 5.00 A is applied?
- (A) $\frac{(2)(55.85)(5.00)}{96500}$
- (B) $\frac{(2)(96500)}{(55.85)(5.00)}$
- (C) $\frac{(55.85)(96500)}{(2)(5.00)}$

(D) $\frac{(2)(55.85)(96500)}{(5.00)}$

Use the following information to answer questions 22-24

10.0 g each of three different gases are present in three glass containers of identical volume, as shown below. The temperature of all three flasks is held constant at 298 K.



22. The container with which gas would have the greatest pressure?

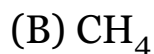
- (A) SO_2
- (B) CH_4
- (C) NCl_3
- (D) All four containers would have the same pressure.

23. Which of the gases would have the greatest density?

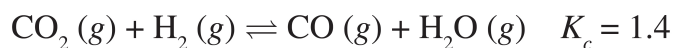
- (A) SO_2
- (B) CH_4
- (C) NCl_3
- (D) All three gases would have the same density.

24. If a small, pinhole-size leak were to be drilled into each container, the container with which gas would experience the fastest pressure decrease?

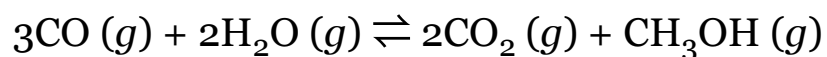
- (A) SO_2



(D) All three containers would decrease pressure at the same rate.



25. Given the above information, what would the equilibrium constant for the below reaction be?

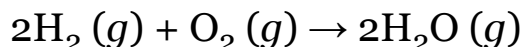


(A) $(2)(1.4)(14.5)$

(B) $\frac{(1.4)(14.5)}{2}$

(C) $\frac{14.5}{(1.4)^2}$

(D) $14.5 - 1.4^2$



26. When 1.0 mole of H_2 is combined with 1.0 mol of O_2 in a sealed flask, the reaction above occurs to completion at a constant temperature. After the reaction, the pressure in the container will have:

(A) Increased by 25%

(B) Increased by 50%

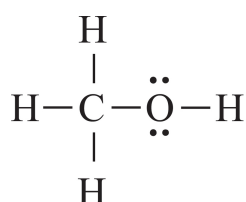
(C) Decreased by 25%

(D) Decreased by 50%

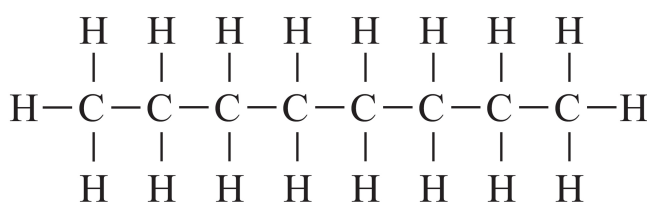
27. A strong acid/strong base titration is completed using an indicator which changes color at the exact equivalence point of

the titration. The protonated form of the indicator is HIn, and the deprotonated form is In⁻. At the equivalence point of the reaction:

- (A) $[\text{HIn}] = [\text{In}^-]$
- (B) $[\text{HIn}] = 1/[\text{In}^-]$
- (C) $[\text{HIn}] = 2[\text{In}^-]$
- (D) $[\text{HIn}] = [\text{In}^-]^2$



Ethanol



Octane

28. The Lewis diagrams for both ethanol and octane are drawn above. Ethanol's boiling point is 78°C, while octane's is 125°C. This is best explained by the fact that:

- (A) Octane has hydrogen bonding, while ethanol does not.
- (B) Octane has a significantly higher molar mass than ethanol.
- (C) Octane's temporary dipoles are stronger than those in ethanol.
- (D) Octane is more symmetrical than ethanol.

29. Which compound, CaCl₂ or CaO, would you expect to have a high melting point? Why?

- (A) CaCl₂, because there are more ions per lattice unit
- (B) CaCl₂, because a chlorine ion is smaller than an oxygen ion

- (C) CaO, because the charge of oxygen ion exceeds that of chlorine ion
- (D) CaO, because the common charges of calcium and oxygen ions are identical in magnitude
30. Even though it is a noble gas, xenon is known to form bonds with other elements. Which element from the options below would xenon most likely be able to bond with?
- (A) Lithium
- (B) Argon
- (C) Fluorine
- (D) Carbon
31. Stock nitric acid, HNO_3 , has a concentration of 15.8 M. The pH of stock nitric acid would fall into which of the following pH ranges?
- (A) Between -2 and -1
- (B) Between -1 and 0
- (C) Between 0 and 1
- (D) Between 1 and 2
32. During gravimetric analysis experiments, collected precipitates are often rinsed with distilled water prior to being collected. What is the purpose of doing so?
- (A) This ensures the precipitation reaction has gone to completion
- (B) It washes away any spectator ions stuck to the precipitate
- (C) The precipitate must be fully hydrated
- (D) Washing the precipitate ensures it has the correct density

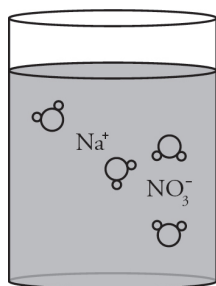


33. The above equilibrium is established in a sealed container. Some neon gas is injected into the container at constant temperature. If the neon does not react with any of the species in the container, what effect will this have on the reaction quotient and the reaction rates for this reaction?
- (A) Q will increase, causing the forward reaction rate to increase
(B) Q will increase, causing the reverse reaction rate to increase
(C) Q will decrease, causing the forward reaction rate to decrease
(D) Q will remain unchanged and the reaction rates will also remain unchanged

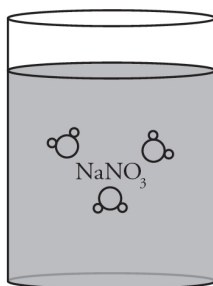
Use the following information to answer question 34-38.

A 10.0 g sample of NaNO_3 is dissolved in 200.0 mL of water with stirring, and the temperature of the water changes from 23.0°C to 18.0°C during the process. Assume the density and the specific heat of the final solution are identical to those of water.

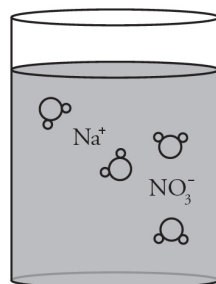
34. Which of the following diagrams correctly shows a particulate representation of the species present in the beaker after the NaNO_3 has dissolved?



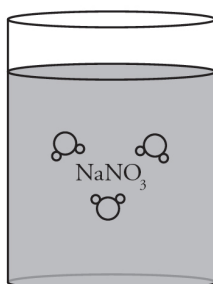
(A)



(C)



(B)



(D)

35. Which of the following statements best describes the enthalpy of solution for NaNO_3 ?

- (A) It will be greater than zero because energy is absorbed from the water
- (B) It will be greater than zero because the magnitude of the ion-dipole attractive forces exceed the lattice energy of NaNO_3
- (C) It will be less than zero because water is a highly polar solvent
- (D) It will be less than zero due to the presence of the polyatomic NO_3^- anion

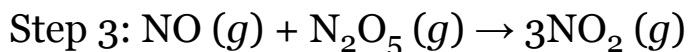
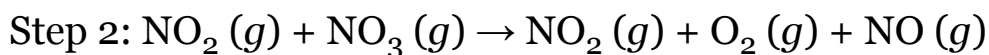
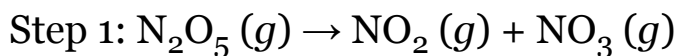
36. During the process, energy is released when:

- (A) The ionic bonds within the NaNO_3 lattice are broken
- (B) The water molecules are spread apart to make room for the dissolved NaNO_3

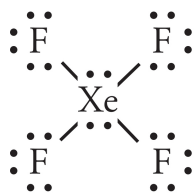
- (C) New attractive forces form between the dissociated ions and the water dipoles
- (D) The covalent bonds within the water molecules are broken
37. Which of the following expressions will correctly calculate the heat change experienced by the water?
- (A) $(200.0 \text{ g})(4.18 \text{ J/g}^\circ\text{C})(-5.0^\circ\text{C})$
- (B) $(210.0 \text{ g})(4.18 \text{ J/g}^\circ\text{C})(-5.0^\circ\text{C})$
- (C) $(10.0 \text{ g})(4.18 \text{ J/g}^\circ\text{C})(-5.0^\circ\text{C})$
- (D) $(200.0 \text{ g})(4.18 \text{ J/g}^\circ\text{C})(5.0^\circ\text{C})$
38. After experimentally determining the enthalpy of solution for NaNO_3 using the gathered data, the magnitude of the experimental value is found to be lower than the accepted value. Which of the following correctly identifies a reason why?
- (A) Some water evaporated while the NaNO_3 was dissolving
- (B) The specific heat of the final solution is actually lower than $4.18 \text{ J/g}^\circ\text{C}$
- (C) The stirring of the solution added significant energy to the system
- (D) Some energy from the dissolution process was lost to the surroundings
39. Red light has a wavelength of 680 nm, and blue light has a wavelength of 470 nm. Which of the following options correctly identifies the relationships between the frequencies and energy levels of red and blue light?

	Frequency	Energy
(A)	Red > Blue	Red > Blue

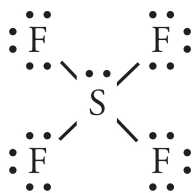
(B)	Red > Blue	Red < Blue
(C)	Red < Blue	Red > Blue
(D)	Red < Blue	Red < Blue



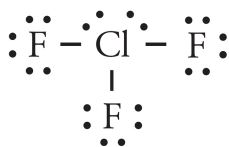
40. The rate law for the above reaction is determined to be $\text{rate} = k[\text{N}_2\text{O}_5]$. Which of the following options correctly identifies the overall order of the reaction as well as the molecularity of the rate-determining step?
- (A) First order, unimolecular
 - (B) Second order, bimolecular
 - (C) Third order, bimolecular
 - (D) Fifth order, unimolecular
41. As temperature decreases, the pH of water increases. This means that:
- (A) The number of H^+ and OH^- ions are increasing
 - (B) Water is more basic at lower temperatures
 - (C) $[\text{H}^+]$ begins to exceed the $[\text{OH}^-]$
 - (D) The auto-ionization of water is an endothermic process
42. The Lewis diagrams for four different compounds are drawn below. In which molecule would the dipole moment be closest to zero?



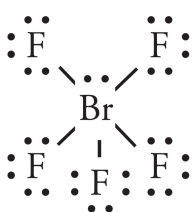
(A)



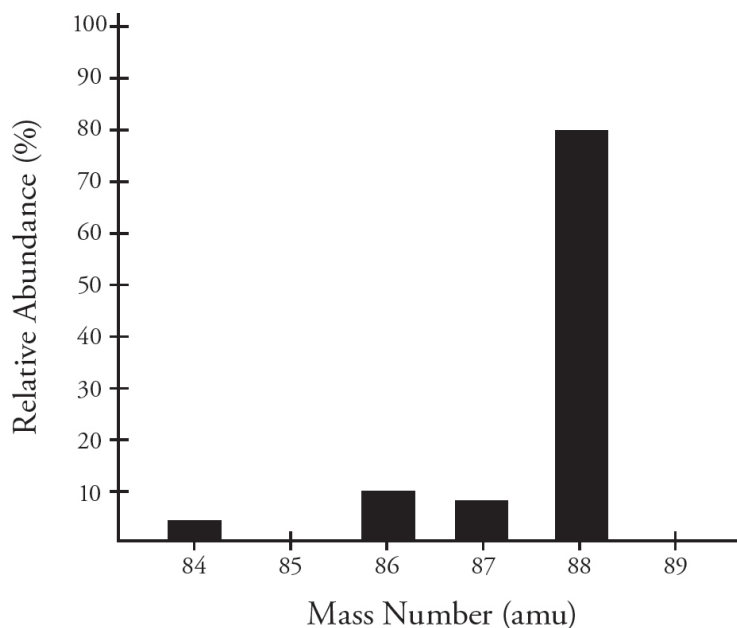
(C)



(B)

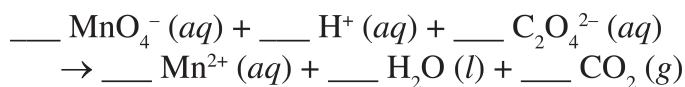


(D)



43. The mass spectrum above shows the distribution of various isotopes of strontium. Based on the data, which of the following conclusions can be drawn?

- (A) Strontium most commonly forms ions with a charge of +2
- (B) Strontium isotopes with a mass of 86 or 87 are very unstable
- (C) The number of protons in a strontium atom nucleus can vary
- (D) The most common isotope of strontium has 50 neutrons

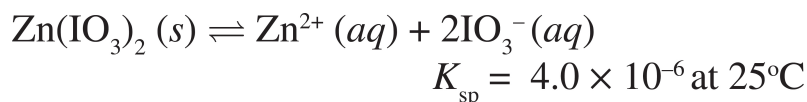


44. When the above oxidation-reduction reaction is completely balanced, what is the coefficient on the MnO_4^- ion?

- (A) 1
- (B) 2
- (C) 3
- (D) 4

Use the following information to answer questions 45-47

Zinc iodate dissociates in water via the following equilibrium process:



45. What is the approximate molar solubility of zinc iodate at 25°C ?

- (A) $1.0 \times 10^{-2} M$
- (B) $2.0 \times 10^{-3} M$
- (C) $4.0 \times 10^{-6} M$
- (D) $1.0 \times 10^{-6} M$

46. Which of the following solutions would zinc iodate be the LEAST soluble in?

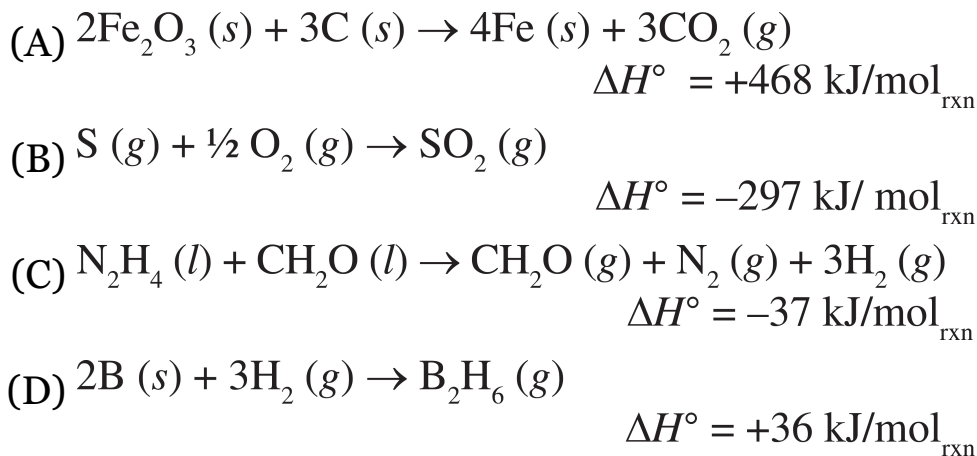
- (A) $1.0 M \text{ BaCl}_2$
- (B) $1.0 M \text{ NaIO}_3$
- (C) $1.0 M \text{ K}_2\text{CO}_3$

(D) Pure water

47. A beaker of saturated zinc iodate is left out overnight. The following morning, some water is found to have evaporated. Assuming the temperature remained constant at 25°C, which of the following options correctly identifies the changes in the $[\text{Zn}^{2+}]$ and the mass of the zinc iodate present in the beaker compared to the previous night?

	$[\text{Zn}^{2+}]$	Mass $\text{Zn}(\text{IO}_3)_2$
(A)	Increase	Decrease
(B)	Increase	No Change
(C)	No Change	Increase
(D)	No Change	Decrease

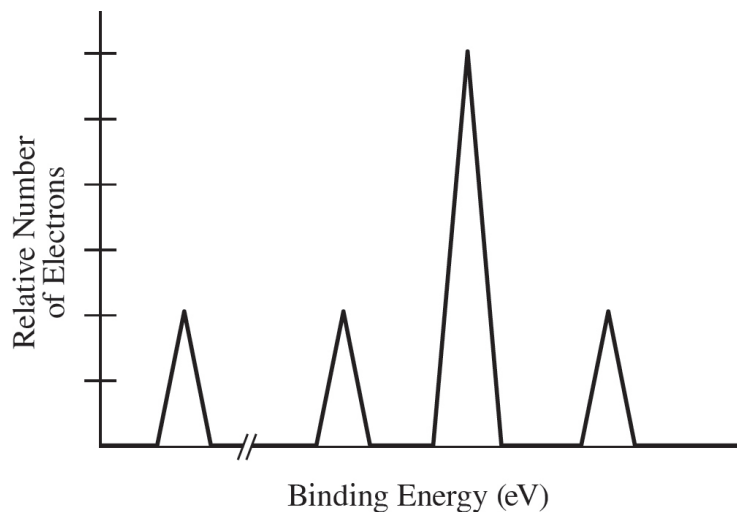
48. Which of the following reactions would be favored at lower temperatures, but not favored at higher temperatures?



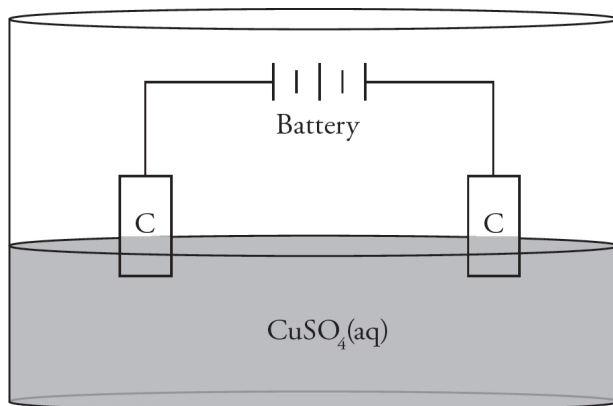
49. Which of the following statements regarding covalent bonds is true?



- (B) Covalent bonds always involve the equal sharing of valence electrons between two atoms
- (C) The breaking of a covalent bond always occurs during a liquid to gas phase change for a covalent substance
- (D) Covalent bonds typically form between atoms that demonstrate metallic properties



50. The photoelectron spectrum of an element is above. Based on the spectrum, what is the charge on the most common ion of the element?
- (A) -2
- (B) -1
- (C) $+1$
- (D) $+2$



51. A beaker is filled with some 1.0 M CuSO_4 , and two carbon electrodes are placed in the beaker with a battery wired between them, as shown above. As current is run through the system, solid copper plates out onto the carbon cathode. Which of the following changes would increase the amount of solid copper that is plated out of solution?

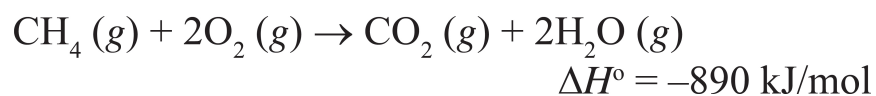
- (A) Replacing the 1.0 M CuSO_4 solution with 1.0 M CuCl_2
- (B) Using a 9.0 V battery instead of a 1.5 V battery
- (C) Decreasing the pH of the solution
- (D) Changing out the carbon electrodes with platinum electrodes



52. 1.0 mole of $\text{NO}_2(g)$ is bubbled through excess water, causing the above reaction to take place. Which of the following statements correctly describes the energy change that will occur during the reaction?

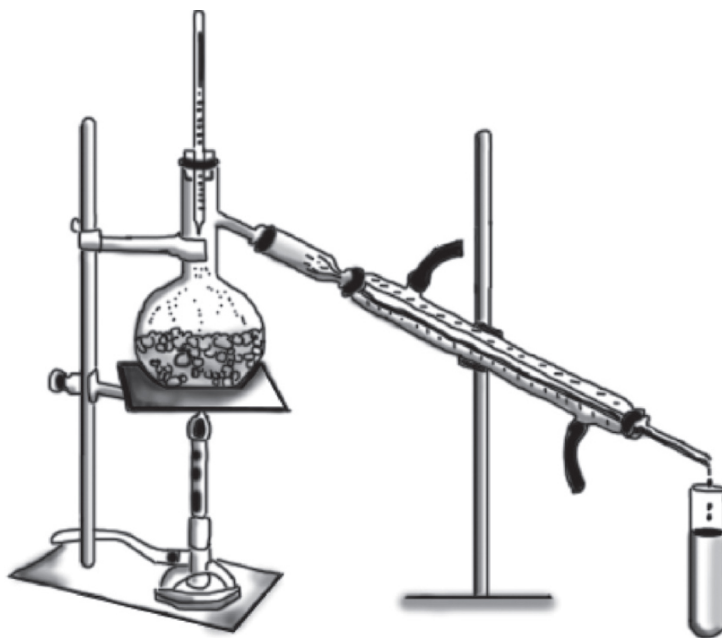
- (A) 135 kJ of energy will be emitted
- (B) 45 kJ of energy will be emitted
- (C) 135 kJ of energy will be absorbed
- (D) 405 kJ of energy will be absorbed

Bond	Energy (kJ/mol)
C-H	413
O=O	498
O-H	467



53. Determine the approximate bond energy of a C=O bond given the above data.

- (A) 400 kJ/mol
- (B) 600 kJ/mol
- (C) 800 kJ/mol
- (D) 1000 kJ/mol



Substance	Vapor Pressure @ 25°C (torr)

H ₂ O	23.8
CH ₃ OH	99.0
C ₆ H ₁₂	78.0

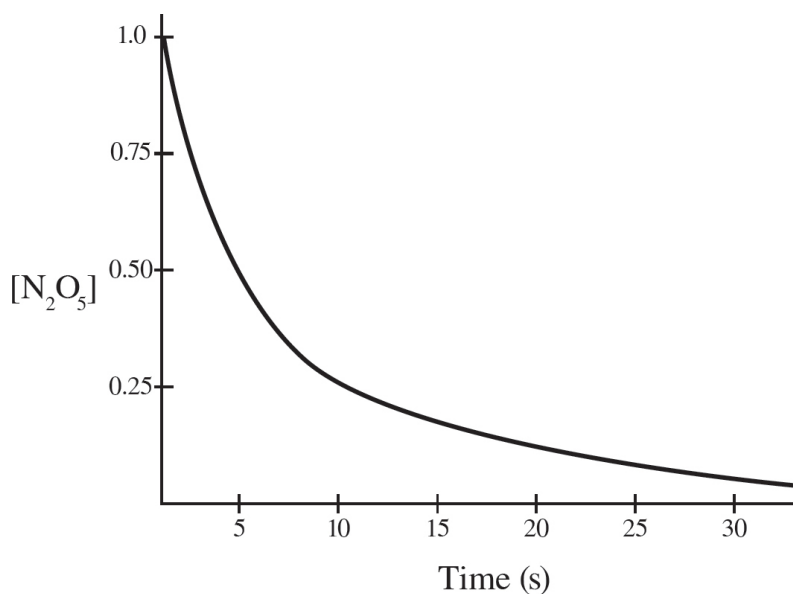
54. Three different liquids are mixed together in a flask, and that flask is then hooked up to a distillation apparatus, as shown above. The liquids are initially at 25°C, and the heat is turned up until the mixture starts to boil. Which liquid would be the first to separate out of the mixture?

(A) H₂O

(B) CH₃OH

(C) C₆H₁₂

(D) Distillation would be an ineffective method of separating the mixture



55. A sample of some N_2O_5 gas is placed in a sealed container and allowed to decompose. The concentration of the N_2O_5 is tracked over time, and the results are plotted on the above graph. Which of the following represent the possible units on the rate constant for this reaction?

- (A) s^{-1}
(B) sM^{-1}
(C) Ms^{-1}
(D) Ms^{-2}



56. The above system is at equilibrium. If it were to be diluted, what would happen to the moles of reactants and the moles product after equilibrium is reestablished?

	Mole Reactants	Moles Product
(A)	Increase	Increase
(B)	Increase	Decrease
(C)	Decrease	Decrease
(D)	No Change	No Change

Use the following information to answer questions 57-60

A beaker contains 100. mL of a 1.0 M solution of benzoic acid, $\text{C}_6\text{H}_5\text{COOH}$, which has a pK_a value of 4.19. 50. mL of Solution X is added to the beaker, creating a buffer with a pH of 4.19.

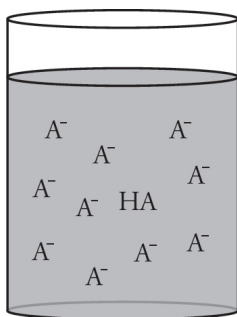
57. Which of the following could be the identity of the Solution X?

- (A) 1.0 *M* NaOH
- (B) 1.0 *M* HCl
- (C) 1.0 *M* NaCl
- (D) 1.0 *M* NH₃

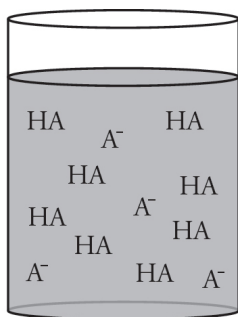
58. If the buffer solution was then diluted with 150. mL of distilled water, which of the following values would be closest to the pH of the diluted buffer solution?

- (A) 2.08
- (B) 4.19
- (C) 5.19
- (D) 6.27

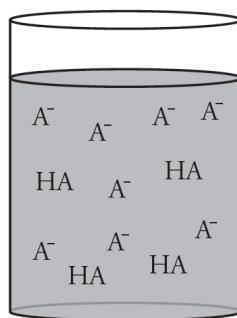
59. In the diagrams below, benzoic acid is represented by HA, and the benzoate ion, C₆H₅COO⁻, is represented by A⁻. Which of the beakers shows the correct ratio of HA:A⁻ in a buffer solution that has a pH of 3.19?



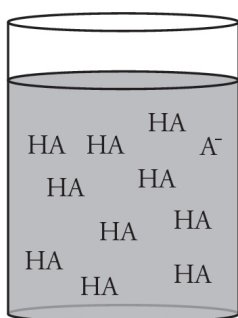
(A)



(C)



(B)



(D)

60. The benzoic acid solution can be made into an effective buffer. Which of the following statements best explain why a HCl solution of identical concentration NOT be part of a buffer?

- (A) The K_a value for benzoic acid is greater than the K_a value for HCl
- (B) HCl has stronger intermolecular forces than C_6H_5COOH
- (C) The Cl^- ion is a less effective conjugate base than $C_6H_5COO^-$
- (D) HCl has far few protons in it than C_6H_5COOH

END OF SECTION I

INFORMATION IN THE TABLE BELOW AND ON THE FOLLOWING PAGES MAY BE USEFUL IN ANSWERING THE QUESTIONS IN THIS SECTION OF THE EXAMINATION.

DO NOT DETACH FROM BOOK.

PERIODIC TABLE OF THE ELEMENTS

PERIODIC TABLE OF THE ELEMENTS																		
1 H 1.008																	2 He 4.00	
3 Li 6.94	4 Be 9.01											5 B 10.81	6 C 12.01	7 N 14.01	8 O 16.00	9 F 19.00	10 Ne 20.18	
11 Na 22.99	12 Mg 24.30											13 Al 26.98	14 Si 28.09	15 P 30.97	16 S 32.06	17 Cl 35.45	18 Ar 39.95	
19 K 39.10	20 Ca 40.08	21 Sc 44.96	22 Ti 47.90	23 V 50.94	24 Cr 52.00	25 Mn 54.94	26 Fe 55.85	27 Co 58.93	28 Ni 58.69	29 Cu 63.55	30 Zn 65.39	31 Ga 69.72	32 Ge 72.59	33 As 74.92	34 Se 78.96	35 Br 79.90	36 Kr 83.80	
37 Rb 85.47	38 Sr 87.62	39 Y 88.91	40 Zr 91.22	41 Nb 92.91	42 Mo 95.94	43 Tc (98)	44 Ru 101.1	45 Rh 102.91	46 Pd 106.42	47 Ag 107.87	48 Cd 112.41	49 In 114.82	50 Sn 118.71	51 Sb 121.75	52 Te 127.60	53 I 126.91	54 Xe 131.29	
55 Cs 132.91	56 Ba 137.33	*La 138.91	Hf 178.49	Ta 180.95	W 183.85	Re 186.21	Os 190.2	Ir 192.2	Pt 195.08	Au 196.97	Hg 200.59	Tl 204.38	Pb 207.2	Bi 208.98	Po (209)	At (210)	Rn (222)	
87 Fr (223)	88 Ra 226.02	†Ac 227.03	Rf (261)	Db (262)	Sg (266)	Bh (264)	Hs (277)	Mt (268)	Ds (271)	Rg (272)								
*Lanthanide Series			58 Ce 140.12	59 Pr 140.91	60 Nd 144.24	61 Pm (145)	62 Sm 150.4	63 Eu 151.97	64 Gd 157.25	65 Tb 158.93	66 Dy 162.50	67 Ho 164.93	68 Er 167.26	69 Tm 168.93	70 Yb 173.04	71 Lu 174.97		
			†Actinide Series		90 Th 232.04	91 Pa 231.04	92 U 238.03	93 Np (237)	94 Pu (244)	95 Am (243)	96 Cm (247)	97 Bk (247)	98 Cf (251)	99 Es (252)	100 Fm (257)	101 Md (258)	102 No (259)	103 Lr (262)

ADVANCED PLACEMENT CHEMISTRY EQUATIONS AND CONSTANTS

Throughout the test the following symbols have the definitions specified unless otherwise noted.

L, mL = liter(s), milliliter(s)
 g = gram(s)
 nm = nanometer(s)
 atm = atmosphere(s)

mm Hg = millimeters of mercury
 J, kJ = joule(s), kilojoule(s)
 V = volt(s)
 mol = mole(s)

ATOMIC STRUCTURE

$$E = h\nu$$

$$c = \lambda\nu$$

E = energy
 ν = frequency
 λ = wavelength

Planck's constant, $h = 6.626 \times 10^{-34}$ J s

Speed of light, $c = 2.998 \times 10^8$ m s⁻¹

Avogadro's number = 6.022×10^{23} mol⁻¹

Electron charge, $e = -1.602 \times 10^{-19}$ coulomb

EQUILIBRIUM

$$K_c = \frac{[C]^c[D]^d}{[A]^a[B]^b}, \text{ where } aA + bB \rightleftharpoons cC + dD$$

$$K_p = \frac{(P_C)^c(P_D)^d}{(P_A)^a(P_B)^b}$$

$$K_a = \frac{[H^+][A^-]}{[HA]}$$

$$K_b = \frac{[OH^-][HB^+]}{[B]}$$

$$K_w = [H^+][OH^-] = 1.0 \times 10^{-14} \text{ at } 25^\circ\text{C}$$

$$= K_a \times K_b$$

$$\text{pH} = -\log[H^+], \text{ pOH} = -\log[OH^-]$$

$$14 = \text{pH} + \text{pOH}$$

$$\text{pH} = \text{p}K_a + \log \frac{[A^-]}{[HA]}$$

$$\text{p}K_a = -\log K_a, \text{ p}K_b = -\log K_b$$

Equilibrium Constants

K_c (molar concentrations)

K_p (gas pressures)

K_a (weak acid)

K_b (weak base)

K_w (water)

KINETICS

$$\ln[A]_t - \ln[A]_0 = -kt$$

$$\frac{1}{[A]_t} - \frac{1}{[A]_0} = kt$$

$$t_{1/2} = \frac{0.693}{k}$$

k = rate constant

t = time

$t_{1/2}$ = half-life

GASES, LIQUIDS, AND SOLUTIONS

$$PV = nRT$$

$$P_A = P_{\text{total}} \times X_A, \text{ where } X_A = \frac{\text{moles A}}{\text{total moles}}$$

$$P_{\text{total}} = P_A + P_B + P_C + \dots$$

$$n = \frac{m}{M}$$

$$K = ^\circ\text{C} + 273$$

$$D = \frac{m}{V}$$

$$KE \text{ per molecule} = \frac{1}{2}mv^2$$

Molarity, M = moles of solute per liter of solution

$$A = abc$$

P = pressure

V = volume

T = temperature

n = number of moles

m = mass

M = molar mass

D = density

KE = kinetic energy

v = velocity

A = absorbance

a = molar absorptivity

b = path length

c = concentration

Gas constant, $R = 8.314 \text{ J mol}^{-1} \text{ K}^{-1}$

$= 0.08206 \text{ L atm mol}^{-1} \text{ K}^{-1}$

$= 62.36 \text{ L torr mol}^{-1} \text{ K}^{-1}$

$1 \text{ atm} = 760 \text{ mm Hg}$

$= 760 \text{ torr}$

STP = 273.15 K and 1.0 atm

Ideal gas at STP = 22.4 L mol^{-1}

THERMOCHEMISTRY/ ELECTROCHEMISTRY

$$q = mc\Delta T$$

$$\Delta S^\circ = \sum S^\circ \text{ products} - \sum S^\circ \text{ reactants}$$

$$\Delta H^\circ = \sum \Delta H_f^\circ \text{ products} - \sum \Delta H_f^\circ \text{ reactants}$$

$$\Delta G^\circ = \sum \Delta G_f^\circ \text{ products} - \sum \Delta G_f^\circ \text{ reactants}$$

$$\Delta G^\circ = \Delta H^\circ - T\Delta S^\circ$$

$$= -RT \ln K$$

$$= -nFE^\circ$$

$$I = \frac{q}{t}$$

q = heat

m = mass

c = specific heat capacity

T = temperature

S° = standard entropy

H° = standard enthalpy

G° = standard free energy

n = number of moles

E° = standard reduction potential

I = current (amperes)

q = charge (coulombs)

t = time (seconds)

Faraday's constant, $F = 96,485 \text{ coulombs per mole of electrons}$

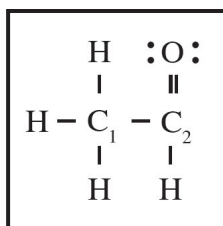
$$1 \text{ volt} = \frac{1 \text{ joule}}{1 \text{ coulomb}}$$

CHEMISTRY
Section II
7 Questions
(Total time—105 minutes)

YOU MAY USE YOUR CALCULATOR FOR THIS SECTION

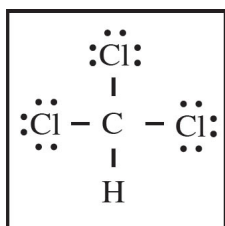
Directions: Questions 1-3 are long free-response questions that require about 23 minutes each to answer and are worth 10 points each. Questions 4-7 are short free-response questions that require about 9 minutes each to answer and are worth 4 points each.

Write your response in the space provided following each question. Examples and equations may be included in your responses where appropriate. For calculations, clearly show the method used and the steps involved in arriving at your answers. You must show your work to receive credit for your answer. Pay attention to significant figures.



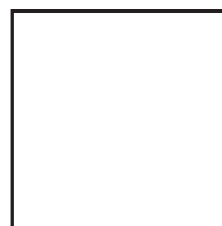
Acetaldehyde, CH_3COH

BP = 293K



Chloroform, CHCl_3

BP = 334K



Methanoic acid, HCOOH

BP = 374K

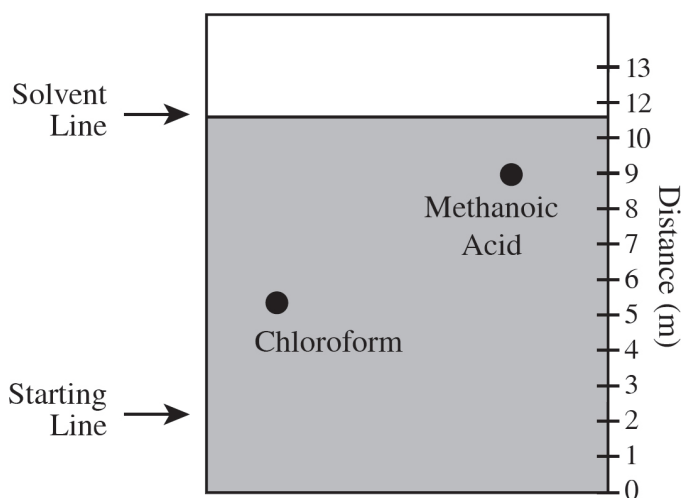
1. The boiling points of three different compounds are listed above, along with their formulas.
 - (a) In the blank box above, draw the Lewis diagram for methanoic acid. There are two carbon-oxygen bonds present, one of which is shorter than the other.
 - (b) (i) What is the hybridization around atom C_2 in the acetaldehyde molecule?
(ii) Chloroform is known to be polar. Draw the location of any positive and negative dipoles on a chloroform

molecule.

- (c) Which of the three molecules would have the greatest degree of polarizability? Justify your answer.

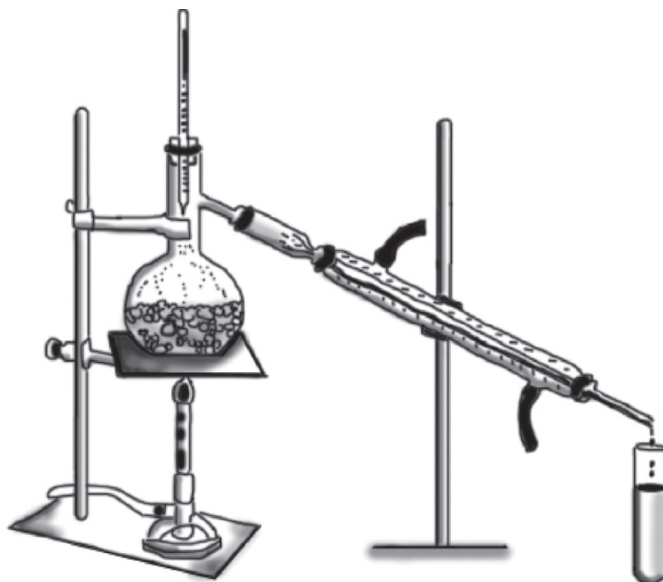
Equal amounts of all three liquids are mixed together inside a beaker.

- (d) A sample of the mixture is placed on a piece of chromatography paper, and the bottom of the paper is immersed in water. After some time passes, the mixture separates. The location of two components of the mixture are charted on the diagram below.



- (i) What is the R_f value for the methanoic acid?
- (ii) On the diagram above draw and label a dot that would be a reasonable estimate of the distance the acetaldehyde would travel.

The beaker containing the mixture is then attached to a distillation apparatus and heated, as shown below.



(e) (i) If the temperature of the beaker is gradually raised, which substance will separate out first? Justify your answer.

(ii) Which of the following temperatures would be the best option in order to create the purest distillate possible? Justify your answer.

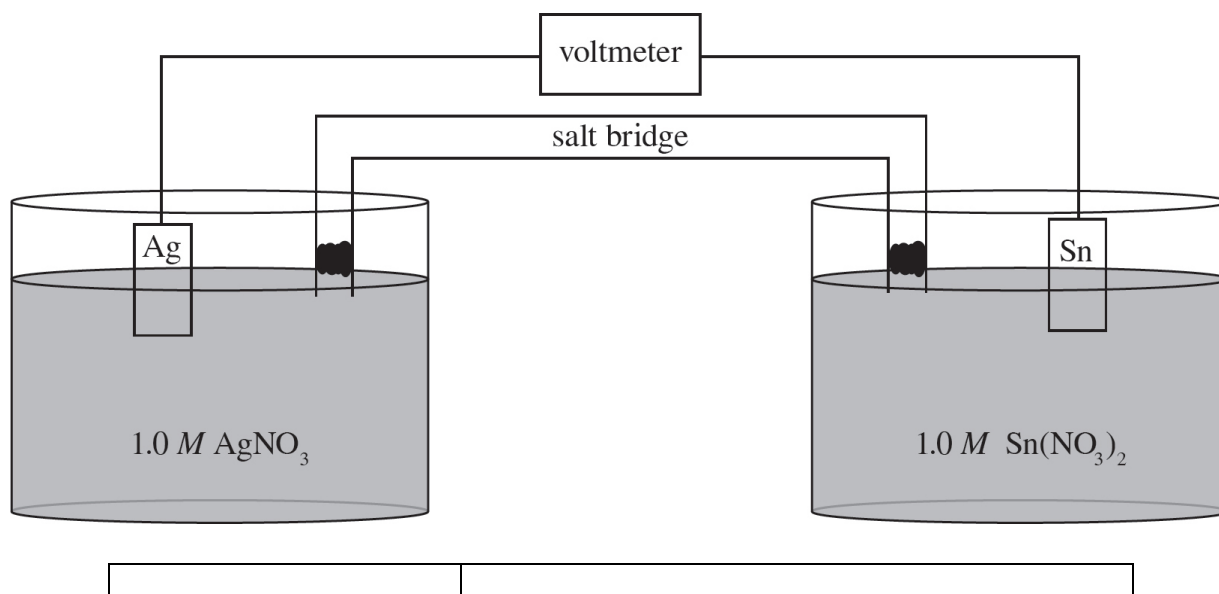
210 K

300 K

330 K

390 K

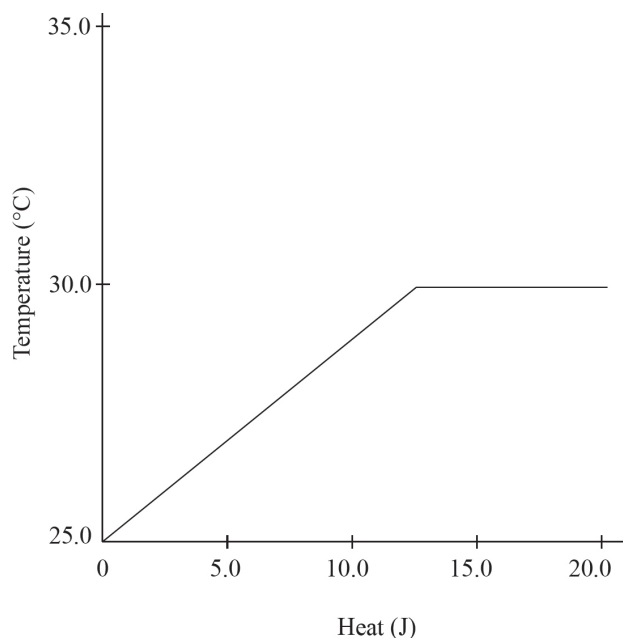
2. A galvanic cell is set up according to the following diagram.



Half-Reaction	Standard Reduction Potential (V)
$\text{Ag}^+ + e^- \rightarrow \text{Ag (s)}$	+0.80
$\text{Sn}^{2+} + 2e^- \rightarrow \text{Sn (s)}$	-0.14

- (a) Write out the balanced net ionic equation that will occur.
- (b) (i) Calculate E°_{cell} .
(ii) Calculate ΔG° for the cell.
- (c) If the solutions in each half-cell were replaced with 2.0 M solutions of AgNO_3 and $\text{Sn}(\text{NO}_3)_2$, how would that effect the cell potential? Justify your answer.
- (d) (i) What is the purpose of the salt bridge?
(ii) Suggest the formula for an aqueous solution that would be a good choice to fill the salt bridge. Justify your answer.
- (e) The beaker of $\text{Sn}(\text{NO}_3)_2$ is disconnected from the cell, and the tin electrode is then connected to an outside source of current. Over the course of 10.0 minutes, 1.65 g of tin plates out onto the electrode. What is the amperage of the current source?

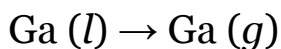
3.



0.10 mol of solid gallium initially at room temperature is heated at a constant rate, and its temperature is tracked, leading to the above graph.

- (a) As heat is added, what is happening to the total entropy of the system? Justify your answer.
- (b) The horizontal portion of the graph indicates a phase change. Explain on a particulate level why the temperature is constant during a phase change.
- (c) (i) Calculate the specific heat capacity of solid gallium in $\text{J g}^{-1} \text{ } ^\circ\text{C}^{-1}$
- (ii) If the specific heat of the solid gallium were greater than what you calculated in (c)(i), how would the slope of the temperature versus heat line change during gallium's solid phase?

The gallium continues to be heated until it fully boils. Assume ideal behavior for the gallium gas.



Substance	(kJ mol ⁻¹)
Ga (l)	5.60
Ga (g)	277.1

(d) (i) Calculate the enthalpy of vaporization for gallium given the above data.

(ii) The enthalpy of vaporization for gallium is greater than its enthalpy of fusion. Explain why in terms of intermolecular forces.

(e) Given your answer to (d)(i) and that $\Delta S^\circ = 113.4 \text{ J mol}^{-1} \text{ K}^{-1}$ for the boiling of gallium, what is the boiling point of the gallium?

(f) After the gallium is fully converted to a gas, it continues to be heated. What would you expect to be true about the velocity distribution of the gaseous gallium atoms as the temperature increases?

4. Sulfurous acid, H_2SO_3 , is a weak diprotic acid.

(a) Write out the two acid-base dissociations which occur when H_2SO_3 is mixed with water.

(b) A 0.50 M solution of H_2SO_3 has a pH of 1.10. Determine K_{a1} for H_2SO_3 .

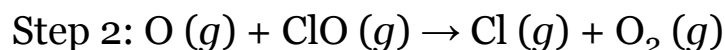
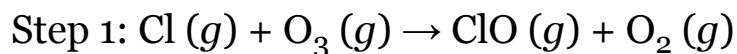
5. Hypobromous acid, HBrO , is a weak monoprotic acid with a K_a value of 2.0×10^{-9} at 25°C.

(a) Write out the equilibrium reaction of hypobromous acid with water, identifying any conjugate acid/base pairs present.

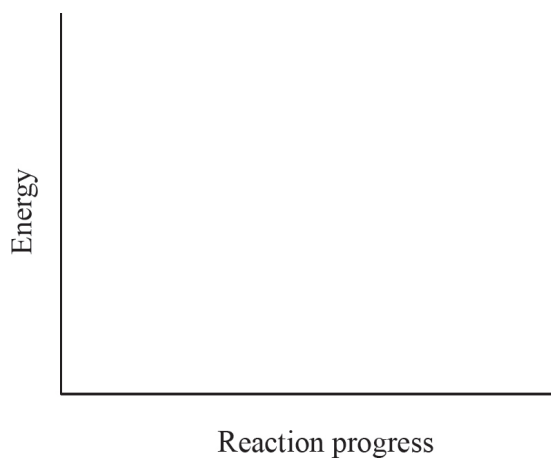
(b) (i) What would be the percent dissociation of a 0.50 M solution of hypobromous acid?

(ii) If the 0.50 *M* solution were diluted, what would happen to the percent dissociation of the HBrO? Why?

6. Chlorofluorocarbons are byproducts of many different processes that are known to be dangerous to the environment as both as a greenhouse gas, as well as an agent for ozone (O₃) depletion. The accepted mechanism for the latter is:



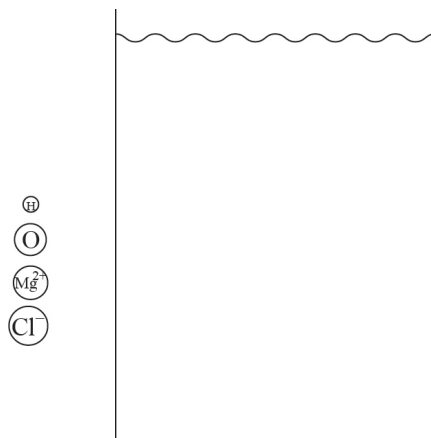
- (a) Write out the full reaction with the above elementary steps, and identify all catalysts and intermediates.
- (b) Describe two ways by which a catalyst can reduce the activation energy of a reaction.
- (c) Both elementary steps in the reaction above are exothermic. On the axes provided, draw a potential reaction mechanism which supports this.



7. A stock solution of 2.0 *M* MgCl₂ is dissolved in water.

- (a) (i) In the beaker below, draw a particulate diagram that represents MgCl₂ dissolved in water. The approximate sizes of each atom/ion are provided for you. Your

diagram should include at least four water molecules, which should be correctly oriented compared to the ions dissolved in solution.



- (ii) Why are the chloride ions from (a)(i) larger than the magnesium ions?
- (b) (i) A student wishes to make up 500 mL of 0.50 M MgCl_2 for an experiment. Explain the best method of doing so, starting with the 2.0 M MgCl_2 solution and utilizing a graduated cylinder and a volumetric flask.
- (ii) What are the concentrations of the Mg^{2+} and Cl^- ions in the new solution?

STOP

END OF EXAM

Practice Test 1:

Answers and Explanations

PRACTICE TEST 1: MULTIPLE-CHOICE ANSWER KEY

- | | | |
|-------|-------|-------|
| 1. A | 21. B | 41. D |
| 2. D | 22. B | 42. A |
| 3. B | 23. D | 43. D |
| 4. A | 24. B | 44. B |
| 5. B | 25. C | 45. A |
| 6. B | 26. C | 46. B |
| 7. C | 27. A | 47. C |
| 8. A | 28. C | 48. B |
| 9. C | 29. C | 49. A |
| 10. D | 30. C | 50. D |
| 11. D | 31. A | 51. B |
| 12. D | 32. B | 52. B |
| 13. B | 33. C | 53. C |
| 14. C | 34. A | 54. B |
| 15. A | 35. A | 55. A |
| 16. A | 36. C | 56. B |
| 17. B | 37. B | 57. A |
| 18. D | 38. D | 58. B |
| 19. A | 39. D | 59. D |
| 20. D | 40. A | 60. C |

SECTION I—MULTIPLE-CHOICE ANSWERS AND EXPLANATIONS

1. A

Molar solubility describes how many moles of a salt will dissolve in one liter of water. Each unit of Na_3PO_4 that dissociates produces three Na^+ ions, which means that there are one-third as many units of Na_3PO_4 as there are of Na^+ .

2. D

The overall dissolution of LiCl is exothermic; this is indicated by the temperature of the water rising. For that to be the case, the energy released during when the ion-dipole attractions form has to exceed the amount of energy required to break the bonds in the LiCl lattice.

3. B

There is a 1:1 mole ratio between the HCl and the CH_3NH_2 , meaning there will be the same number of moles of each present at the equivalence point. The equivalence point is located at 15.0 mL of HCl added. $(1.0\text{ M})(0.0150\text{ L}) = 0.0150\text{ mol HCl} = 0.0150\text{ mol CH}_3\text{NH}_2$. Finally, divide that value by the volume of the CH_3NH_2 , $0.0150\text{ mol CH}_3\text{NH}_2 / 0.020\text{ L} = 0.75\text{ M CH}_3\text{NH}_2$.

4. A

At the half-equivalence point (7.5 mL) of the titration, $\text{p}K_{\text{b}}$ of methylamine = pOH of the solution. To determine the pOH , you simply have to see what the pH of the solution is

at that half-equivalence point and use $\text{pH} + \text{pOH} = 14$. The pH at the half-equivalence is 10.5. So, $14 - 10.5 = 3.5$.

5. **B**

A buffer solution is one that resists change in pH due to similar amounts of the base (CH_3NH_2) and its conjugate acid (CH_3NH_3^+). That occurs after the initial pH change, but prior to the equivalence region. The pH is also stable again starting at 16.5 mL, but that is due to the presence of the excess strong acid, and that does not create a buffer region.

6. **B**

Sodium hydroxide is a strong base which dissociates completely in solution. Thus, it would initially have a higher pH (be more basic) than any weak base of an identical concentration. At equivalence of a strong acid/strong base titration, water is the only acid or base present, causing the solution to have a neutral pH of 7. Finally, in the post-equivalence region of the graph, the pH is driven by the excess H^+ from the HCl, and that would not change in the new titration.

7. **C**

The H—C bond is a single bond with a bond order of one. The C—O bonds display resonance, and the average bond order between them is $(1 + 2)/2 = 1.5$.

8. **A**

Diluting an aqueous system at equilibrium will cause a shift to the side with more aqueous species. In this case, that

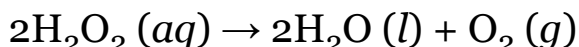
entails a shift to the left, which is what happens when the reverse reaction rate increases.

9. **C**

In order for the reaction to occur, one of the oxygen atoms from NO_2 must have collided directly with the carbon atom in CO . This will allow for the bonds in NO_2 to be broken, while a new C—O bond is also created.

10. **D**

Given that both products are gases at the same temperature and pressure, their volumes are directly proportional with how many moles of gas are present. So, 1.5 times as much CO_2 was created as H_2O . However, you are interested in the C:H ratio, and there are two hydrogen atoms in each molecule of H_2O . The C:H ratio then becomes 1.5:2.0, which is the same as 3:4.



11. **D**

In hydrogen peroxide, oxygen takes on an oxidation state of -1 . Oxygen's oxidation state is -2 in water (a reduction) and 0 in oxygen gas (an oxidation). The oxidation state on hydrogen is a $+1$ in both compounds it appears in.

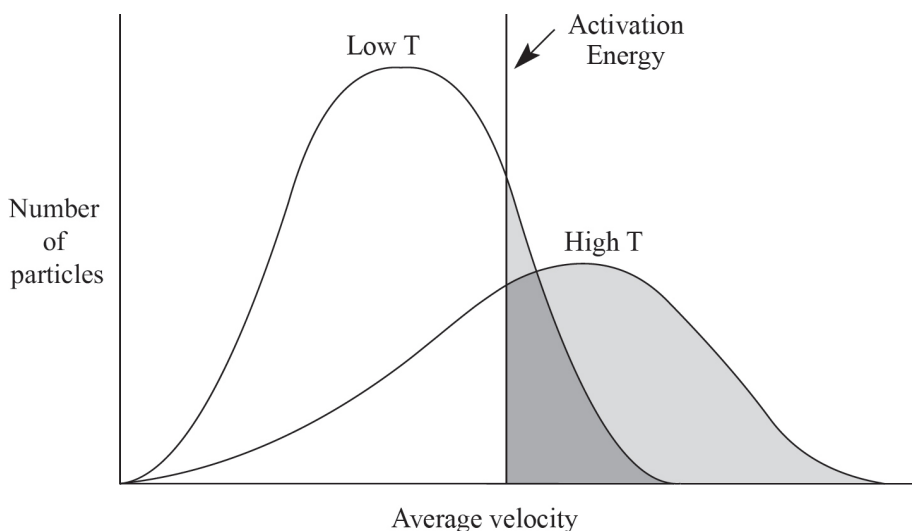
12. **D**

Gases deviate from ideal behavior due to the strength of their IMFs. All four diatomic gases listed are non-polar, so the only difference in IMF strength arises from the polarizability of their electron clouds. Fluorine has the most electrons, and thus, experiences the strongest London

Dispersion Forces and the greatest deviation from ideal behavior.

13. **B**

With increasing temperature, the Maxwell-Boltzmann distribution curve shifts rightward and downward. A larger percentage of the curve will be past the activation energy barrier, as shown below, where the shaded areas under each curve are the particles that have enough energy to react.



(A) is a good distractor, however, just because the average kinetic energy present in the sample increases does not mean the velocity of ALL of the molecules increases (as shown in the distribution curves above).

14. **C**

Phosphorus forms ions with a -3 charge (P^{3-}). Adding valence electrons increase the repulsion between the existing electrons, which increases the size of the ion. Zinc, on the other hand, loses electrons when forming ions, decreasing repulsion and reducing size.

15. **A**

Phosphorus has electrons in both the $3s$ and $3p$ sublevels. These sublevels are located on the third energy level and are further away from the nucleus than the electrons in the first two energy levels, which are the only ones neon has electrons in.

16. **A**

When transition metals form ions, they lose their outermost s -electrons first. Additionally, when filling the d -orbitals, you always need to subtract one from the period the element is found in to determine the principal energy level of its d -orbital.

17. **B**

Both molecules have four electron groups around the central atom, so their base geometry is tetrahedral, which has bond angles of 109.5° . However, the PCl_3 has a lone pair as well, which exerts a slightly greater repulsive force than a bonded pair, bringing the bond angles in that molecules to slightly less than the ideal 109.5° .

18. **D**

Stirring will only increase the rate of a reaction that is heterogeneous in nature. In this example, all reactants and products are in the liquid or aqueous phase, meaning the inherent speed of the molecules and ions is so great that any energy added via stirring would be negligible.

19. **A**

Vapor pressure arises from the fact that molecules on the surface of a liquid may have enough kinetic energy to

escape the intermolecular forces holding them together in liquid form. As temperature increases, the average overall kinetic energy of the molecules increases, meaning more of them will have enough energy to escape the surface of the liquid.

20. **D**

Enthalpy of formation involves the formation of one mole of the substance from elements in their standard states. Both hydrogen and oxygen are diatomic elements in their standard state. Note that halves are considered acceptable when balancing diatomic elements.

21. **B**

The best way to look at this is as an exercise in unit analysis. That means you need to start with the initial unit (grams), and find conversions to get the desired unit (seconds). First, go from grams to moles using the molar mass of the compound. It takes 2 moles of electrons to convert one mole of Fe^{2+} to Fe (s). There are 96,500 Coulombs of charge in one mole of electrons (Faraday's constant). Finally, 5.0 Amps = 5.0 Coulombs per second. Putting it all together:

$$1.00 \text{ g Fe} \times \frac{1 \text{ mol Fe}}{55.85 \text{ g Fe}} \times \frac{2 \text{ mol } e^-}{1 \text{ mol Fe}} \times \frac{96,500 \text{ C}}{1 \text{ mol } e^-} \times \frac{1.0\text{s}}{5.00 \text{ C}} = \text{Time in seconds}$$

22. **B**

Pressure is directly related to the number of moles of a gas. CH_4 has the lowest molar mass out of the three gases, and therefore 10.0 grams of it represents the greatest number of moles, leading to the greatest pressure.

23. **D**

Density is simply mass divided by volume. The mass and volume of each gas is identical, therefore, their densities are identical.

24. **B**

The rate at which a gas effuses is inversely proportional with its molar mass. A gas with a lower molar mass (in this case, CH_4), will have molecules moving with a higher average velocity, meaning they are going to hit that hole more often and will be more likely to escape.

25. **C**

To get the targeted equation, you have to flip the top one and multiple it by two before combining it with the bottom one. When manipulating equilibrium constants, flipping a reaction forces you to take the reciprocal of the original K , and doubling all coefficients leads to K being squared. So for $2\text{CO} (g) + 2\text{H}_2\text{O} (g) \rightleftharpoons 2\text{CO}_2 (g) + 2\text{H}_2 (g)$, $K_c = (1/1.4)^2$. After that, when you add the two equations together, to get the K value for their sum you have to multiple the two K values together.

26. **C**

First, the limiting reactant is clearly the H_2 , as twice as much of it is needed for the reaction to go to completion, yet there is not twice of much as it present. With that in mind, the ICE chart is as follows:

	2H_2	O_2	$2\text{H}_2\text{O}$
Initial	1.0	1.0	0
Change	$-2x$	$-x$	$+2x$
Ending	0		

A quick solve tells you x is 0.5, meaning that at the end of the reaction there are 0.5 moles of oxygen and 1.0 moles of H_2O . That is 1.5 total moles of gas left at the end, a 25% decrease in moles (and thus, pressure) from the 2.0 moles of gas that was present initially.

27. **A**

An indicator changes color when the concentration of the deprotonated form exceeds the concentration of its protonated form. At the exact point of the color change, the concentrations of both the protonated and deprotonated forms will be equal.

28. **C**

Ethanol displays hydrogen bonding, which is the strongest type of intermolecular force. However, London dispersion forces are based on the overall polarizability of a molecule's electron cloud, which in turn is based on the number of electrons that molecule has. Octane has significantly more electrons than ethanol, and the dispersion forces caused by that overwhelm even the hydrogen bonds in ethanol, causing octane to have a higher boiling point.

29. **C**

When considering melting points for ionic compounds, a greater lattice energy means a greater melting point. Lattice energy is calculated using Coulomb's Law, which tells us that both greater charge and smaller size lead to higher energy. The oxygen anion has a charge of -2 , while chlorine's anion charge is only -1 . Note that oxygen is also smaller than chlorine, so that further reinforces the greater melting point of CaO .

30. **C**

Noble gases don't usually form compounds because they have full outermost energy levels and are very stable. However, the larger noble gases (xenon and krypton) have valence electrons far enough away from the nucleus that they are able to share those with highly electronegative elements such as fluorine or oxygen.

31. **A**

$\text{pH} = -\log[\text{H}^+]$. A 10 *M* solution of a strong acid such as nitric would have a pH of $-\log(10) = -1$. As 15.8 *M* is greater than 10 *M*, the pH of the 15.8 *M* solution must be more acidic (lower).

32. **B**

Even though the spectator ions do not take part in the actual precipitation reaction, they can still “stick” to the surface of the precipitate itself. Rinsing the precipitate with water will carry those ions away as the dipoles in the water attract the free spectator ions.

33. **C**

$Q = (\text{P}_{\text{SO}_2})^2(\text{P}_{\text{O}_2})/(\text{P}_{\text{SO}_3})^2$. Injecting neon into the tank will reduce the mole fraction, and thus the partial pressure, of all the gases. Reducing the partial pressures will decrease the numerator more than the denominator due to the ratios in *Q*. This, in turn, causes the value for *Q* to decrease, necessitating a shift to the right in order to re-establish equilibrium.

34. **A**

The Na^+ cations will be attracted to the negative (oxygen) dipoles on the water molecules, and the NO_3^- anions will

be attracted to the positive (hydrogen) dipoles.

35. **A**

The temperature drops, which means the reaction absorbed energy from the water. This means the reaction itself is endothermic, giving it a positive value for ΔH° .

36. **C**

The formation of the ion-dipole attractions is an exothermic process during which energy is released.

37. **B**

The equation here is $q = mc\Delta T$, where m is the combined mass of the water and the dissolved salt. The density of pure water is 1.0 g/mL, so 200.0 mL of water has a mass of 200.0 g, and the total mass of the solution is thus 210 g.

38. **D**

If some of the temperature change of the water was due to heat flowing into the surroundings instead of into the dissolution itself, that would cause the overall magnitude of heat change, and thus the enthalpy of solution, to be artificially low.

39. **D**

Via $c = \nu\lambda$, and given that c is a constant (3.0×10^8 m/s), the higher the wavelength, the lower the frequency. Via $E = h\nu$, with h being Planck's constant, a higher frequency means more energy.

40. **A**

There is only one reactant in the rate law, and it is first order. This means the reaction is first order. The rate-determining step is the one in which the coefficient on the reactant matches its order. That would be the first step, and there is only a single reactant molecule in that step, making it unimolecular.

41. **D**

The auto-ionization of water is represented by $\text{H}_2\text{O} (l) \rightleftharpoons \text{H}^+ + \text{OH}^-$. An increase in pH means a decrease in $[\text{H}^+]$, and as pure water is always neutral, a corresponding decrease in $[\text{OH}^-]$ number of OH^- ions. This would only occur if the reaction experiences a permanent shift to the left, and that would only occur during a temperature decrease if the reaction were endothermic.

42. **A**

Molecules that are square planar, like XeF_4 , are symmetrical and thus, nonpolar.

43. **D**

The most common isotope of strontium has an atomic mass of 88, and a strontium nucleus has 38 protons. As atomic mass = protons + neutrons, that means the most common isotope of strontium must have 50 neutrons.

44. **B**

The manganese goes from an oxidation state of +7 in MnO_4^- to +2 in Mn^{2+} , necessitating a gain of 5 electrons. The carbon goes from an oxidation state of +3 in $\text{C}_2\text{O}_4^{2-}$ to +4 in CO_2 , and in the balanced equation that would happen twice (i.e., a 2 coefficient is needed in front of the CO_2 to

balance it), necessitating a loss of 2 electrons. To balance the charge, the manganese half-reaction must be multiplied by 2 while the carbon half-reaction is multiplied by 5.

45. **A**

$$K_{\text{sp}} = [\text{Zn}^{2+}][\text{IO}_3^-]^2$$

$$4.0 \times 10^{-6} = (x)(2x)^2$$

$$4.0 \times 10^{-6} = 4x^3$$

$$1.0 \times 10^{-6} = x^3$$

$$x = 1.0 \times 10^{-2} M$$

46. **B**

Due to the common ion effect, zinc iodate would be least soluble in a solution with which is shared an ion. Of the options, only NaIO_3 shares an ion (IO_3^-) with $\text{Zn}(\text{IO}_3)_2$.

47. **C**

In a saturated solution, the concentration of each component ion will always remain constant. For that to happen if water evaporates, some ions must “fall” out of solution, increase the total amount of precipitate.

48. **B**

Given that $\Delta G = \Delta H - T\Delta S$, for a reaction to only be favored at lower temperatures ΔH must be negative, and the $-T\Delta S$ term must be positive (and smaller than ΔH). For the $-T\Delta S$ term to be positive, ΔS itself must be negative, which is the case with reaction (B), as the disorder decreases when 1.5 moles of gaseous reactants turn into 1.0 mole of gaseous products.

49. **A**

The reason that atoms bond is to minimize the potential energy in a system.

50. **D**

The PES represents an electron configuration of $1s^2 2s^2 2p^6 3s^2$ that of magnesium. When forming an ion, the most stable pathway is to lose the two electrons in the third energy level, giving the ion a charge of +2.

51. **B**

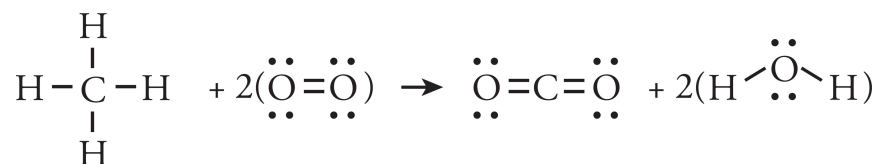
Using a battery with a larger voltage would lead to a larger current, which in turn would lead to a larger mass of copper being plated out of solution.

52. **B**

$$1.0 \text{ mol NO}_2 \times \frac{1 \text{ mol}_{\text{rxn}}}{3 \text{ mol NO}_2} \times \frac{-135 \text{ kJ}}{1 \text{ mol}_{\text{rxn}}} = -45 \text{ kJ} = -45 \text{ kJ}$$

(the negative sign means energy is released).

53. **C**



$$\begin{array}{c} \text{C-H} \\ 4(413) \end{array} + \begin{array}{c} \text{O=O} \\ 2(498) \end{array} - 2x - \begin{array}{c} \text{C=O} \quad \text{O-H} \\ 4(467) \end{array} = -890 \text{ kJ/mol}$$

$$x = +800 \text{ kJ/mol}$$

54. **B**

Vapor pressure is inversely proportional with IMF strength, so the substance with the greatest vapor pressure (in this case, CH_3OH) would have the weakest IMF strength. That also means it has the lowest boiling point, and would boil first.

55. **A**

The graph shows a constant half-life—that is, the concentration of the N_2O_5 decreases by half every 5 seconds. The rate law is thus $\text{rate} = k[\text{N}_2\text{O}_5]$, dimensional analysis gets us to the units on k : $\text{M/s} = k(\text{M})$. $k = \text{s}^{-1}$.

56. **B**

The reaction quotient is $Q = [\text{Cu}(\text{NH}_3)_4^{2+}]/[\text{Cu}^+][\text{NH}_3]^4$. Diluting the equilibrium will decrease the concentration of all ions by the same amount, but that will have a larger effect on the denominator in Q because there are more ions present. This, in turn, means Q will increase, causing a shift to the left, increasing the amount of reactants and decreasing the amount of products.

57. **A**

Adding NaOH to the beaker would cause the following reaction to occur: $\text{C}_6\text{H}_5\text{COOH} + \text{OH}^- \rightleftharpoons \text{C}_6\text{H}_5\text{COO}^- + \text{H}_2\text{O}$. This creates sufficient conjugate base ($\text{C}_6\text{H}_5\text{COO}^-$) to create a buffer system with the benzoic acid.

58. **B**

Diluting a buffer does not change its pH appreciatively, as the concentration of both the acid and its conjugate change equally, which cancels out in Henderson-Hasselbalch. The

pH will become a little bit closer to 7 (the pH of pure water), but not significantly so.

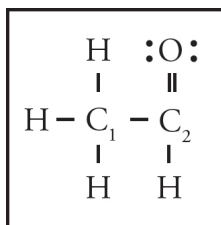
59. **D**

For the pH to decrease by 1 from the pK_a value, via Henderson-Hasselbalch ($pH = pK_a + \log \frac{[C_6H_5COO^-]}{[C_6H_5COOH]}$) the concentration of the acid must be 10 times greater than the concentration of the conjugate base.

60. **C**

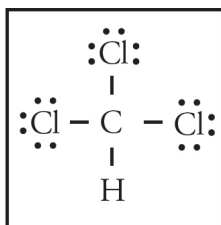
The conjugate bases of strong acids are ineffective, due to the fact that strong acids dissociate fully in solution. Without an effective conjugate base, a buffer cannot be formed.

SECTION II—FREE-RESPONSE ANSWERS AND EXPLANATIONS



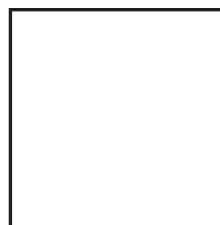
Acetaldehyde, CH_3COH

BP = 293K



Chloroform, CHCl_3

BP = 334K

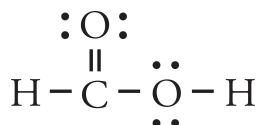


Methanoic acid, HCOOH

BP = 374K

1. The boiling points of three different compounds are listed above, along with their formulas.

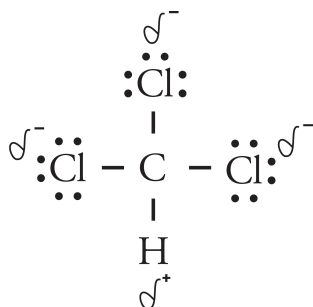
- (a) In the blank box above, draw the Lewis diagram for methanoic acid. There are two carbon-oxygen bonds present, one of which is shorter than the other.



- (b) (i) What is the hybridization around atom C_2 in the acetaldehyde molecule?

Three charge clouds means the hybridization is sp^2 .

- (ii) Chloroform is known to be polar. Draw the location of any positive and negative dipoles on a chloroform molecule.

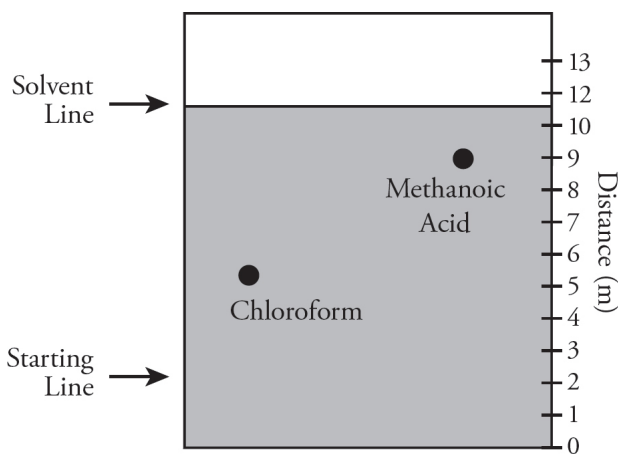


- (c) Which of the three molecules would have the strongest London dispersion forces? Justify your answer.

The polarizability is based on the number of electrons each molecule has. More electrons leads to stronger temporary dipoles, and of the three molecules chloroform has the most electrons and thus the most polarizable.

Equal amounts of all three liquids are mixed together inside a beaker.

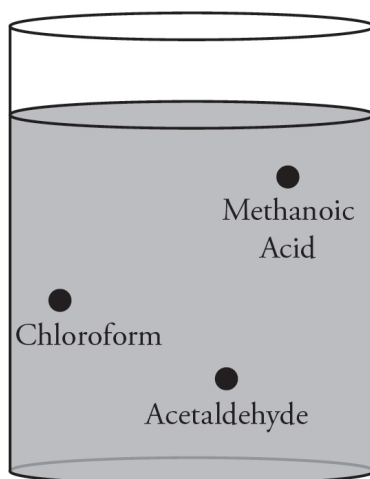
- (d) A sample of the mixture is placed on a piece of chromatography paper, and the bottom of the paper is immersed in water. After some time passes, the mixture separates. The location of two components of the mixture are charted on the diagram below.



- (i) What is the R_f value for the methanoic acid?

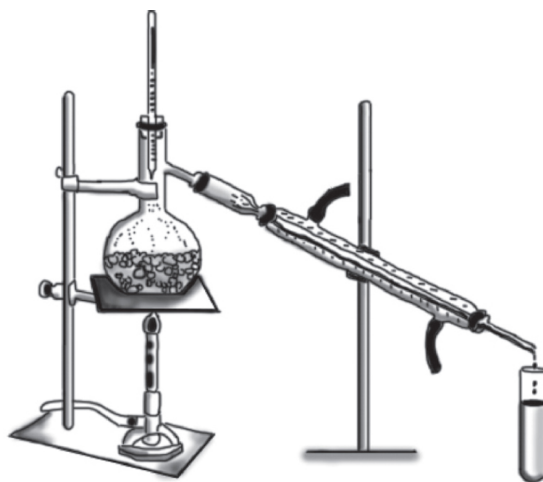
$$R_f = \frac{\text{distance methanoic acid}}{\text{distance solvent}} = \frac{9 \text{ cm}}{11 \text{ cm}} = 0.81$$

- (ii) On the diagram above draw and label a dot that would be a reasonable estimate of the distance the acetaldehyde would travel.



The beaker containing the mixture is then attached to a distillation apparatus and heated, as shown below.

The beaker containing the mixture is then attached to a distillation apparatus and heated, as shown below.



- (e) (i) If the temperature of the beaker is gradually raised, which substance will separate out first? Justify your

answer.

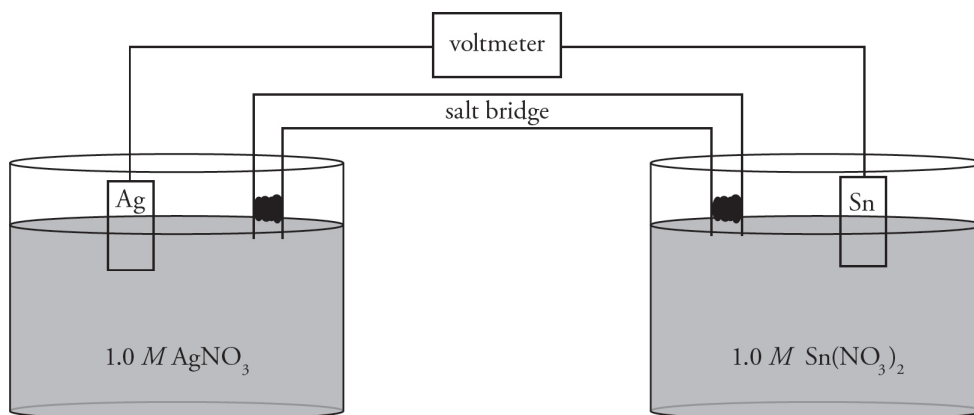
The acetaldehyde is the substance with the lowest boiling point and thus will boil first.

- (ii) Which of the following temperatures would be the best option in order to create the purest distillate possible? Justify your answer.

210 K 300 K 330 K 390 K

300 K is above the boiling point of acetaldehyde and will cause it to boil, but below the boiling points of the other substances, meaning they will not boil. 330 K also fits those guidelines, however, the vapor pressure of the other substances would increase with increasing temperature, and more of the methanoic acid and the chloroform would end up mixed in with the acetaldehyde distillate.

2. A galvanic cell is set up according to the following diagram.



Half-Reaction	Standard Reduction Potential (V)
$\text{Ag}^+ + e^- \rightarrow \text{Ag (s)}$	+0.80
$\text{Sn}^{2+} + 2e^- \rightarrow \text{Sn}$	-0.14

(s)	
-----	--

- (a) Write out the balanced net ionic equation that will occur.

Ag^+ has a higher Standard Reduction Potential than Sn^{2+} , so Ag^+ will be reduced ($\text{Ag}^+ + e^- \rightarrow \text{Ag (s)}$) and Sn (s) will be oxidized ($\text{Sn (s)} \rightarrow \text{Sn}^{2+}$). The reduction reaction must be multiplied by 2 to balance the charges when the half-reactions are combined, so the full net ionic equation becomes $2\text{Ag}^+ + \text{Sn (s)} \rightarrow \text{Sn}^{2+} + 2\text{Ag (s)}$

- (b) (i) Calculate E°_{cell} .

$$+0.80 \text{ V} - (-0.14 \text{ V}) = +0.94 \text{ V}$$

- (ii) Calculate ΔG° for the cell.

$\Delta G^\circ = -nFE$. The number of moles of electrons transferred in this reaction is 2, so $\Delta G^\circ = -(2 \text{ mol } e^-)(96500 \text{ C/mol } e^-)(+0.94 \text{ V}) = -180,000 \text{ J/mol}$ or -180 kJ/mol

- (c) If the solutions in each half-cell were replaced with 2.0 M solutions of AgNO_3 and $\text{Sn(NO}_3)_2$, how would that effect the cell potential? Justify your answer.

Given that $Q = [\text{Ag}^+]^2/[\text{Sn}^{2+}]$, doubling the concentration of each ion would lead to an overall increase in Q . As the cell is favored, the equilibrium constant is greater than 1, so an increase in Q brings the cell closer to equilibrium, when the voltage would be zero. Thus, the cell potential would decrease if 2.0 M solutions were used.

- (d) (i) What is the purpose of the salt bridge?

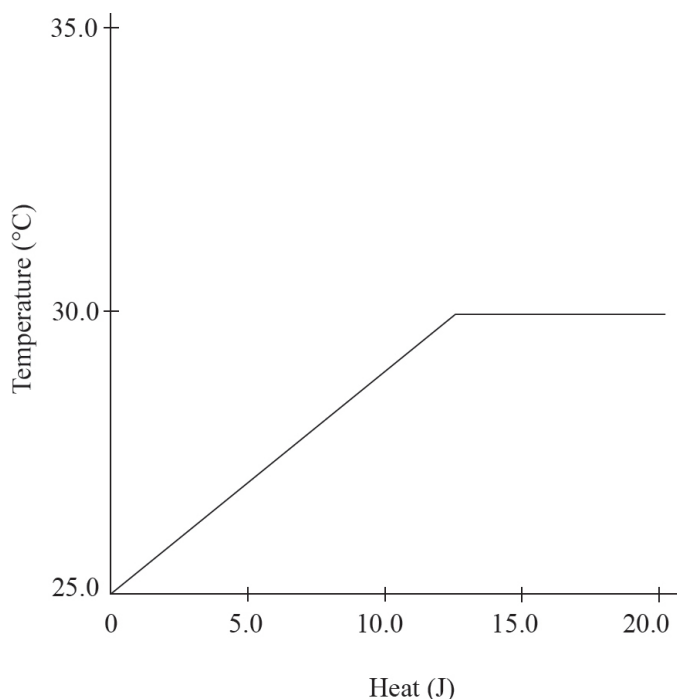
The salt bridge allows ions to flow to each half-cell in order to maintain the electrical neutrality of the cell.

- (ii) Suggest the formula for an aqueous solution that would be a good choice to fill the salt bridge. Justify your answer.

Any aqueous solution that prevents the formation of a precipitate with the ions present in either half-cell would be ideal. As neither alkali metals nor nitrates can form precipitates, a solution of NaNO_3 or KNO_3 would work, as would any other alkali nitrate or ammonium nitrate.

- (e) The beaker of $\text{Sn}(\text{NO}_3)_2$ is disconnected from the cell, and the tin electrode is then connected to an outside source of current. Over the course of 10.0 minutes, 1.65 g of tin plates out onto the electrode. What is the amperage of the current source?

$$1.65 \text{ g Sn} \times \frac{1 \text{ mol Sn}}{118.71 \text{ g}} \times \frac{2 \text{ mol } e^-}{1 \text{ mol Sn}} \times \frac{96500 \text{ C}}{1 \text{ mol } e^-} = \frac{2680 \text{ C}}{600 \text{ s}} = 4.47 \text{ A}$$



0.10 mol of pure gallium initially at room temperature is heated at a constant rate, and its temperature is tracked, leading to the graph.

- (a) As heat is added, what is happening to the total entropy of the system? Justify your answer.

The entropy of the system is increasing as it becomes more disordered. Even in a solid phase, heating a substance causes the vibration of the particles to increase, meaning a greater level of disorder in the system.

- (b) The horizontal portion of the graph indicates a phase change. Explain on a particulate level why the temperature is constant during a phase change.

During a phase change from solid to liquid, the heat is no longer causing the atoms of the gallium to change speed. Instead, the heat goes into weakening the attractions

between the gallium atoms in order to enact the phase change.

- (c) (i) Calculate the specific heat capacity of solid gallium in $\text{J g}^{-1} \text{ } ^\circ\text{C}^{-1}$

From the graph, you can see that it takes approximately 12.0 J of heat to raise the temperature of the gallium by 5.0°C . Additionally, $0.10 \text{ mol Ga} \times 69.7 \text{ g/mol} = 6.97 \text{ g}$ of gallium.

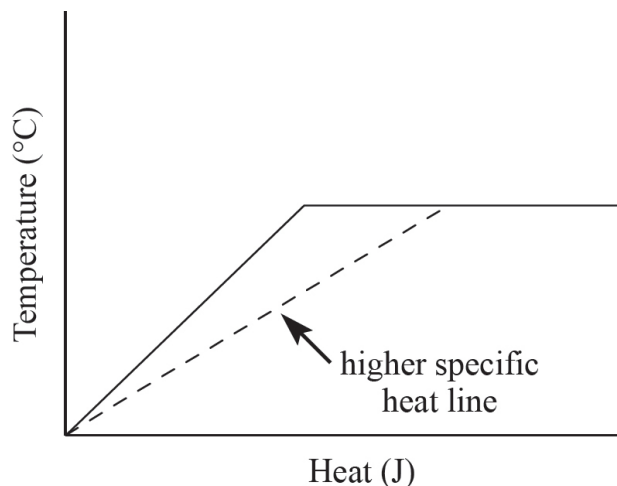
$$q = mc\Delta T$$

$$12.0 \text{ J} = (6.97 \text{ g})(c)(5.0^\circ\text{C})$$

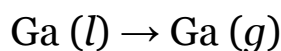
$$c = 0.344 \text{ J g}^{-1} \text{ } ^\circ\text{C}^{-1}$$

- (ii) If the specific heat of the solid gallium were greater than what you calculated in (c)(i), how would the slope of the temperature versus heat line change during gallium's solid phase?

If the specific heat were actually greater, it would take more energy to raise the temperature of the gallium, meaning the slope of the line would be smaller, as shown below. Note that the melting point of the gallium would be unchanged, it would just take more heat to raise the temperature of the solid gallium to that point.



The gallium continues to be heated until it fully boils. Assume ideal behavior for the gallium gas.



Substance	$\Delta(\text{kJ mol}^{-1})$
Ga (l)	5.60
Ga (g)	277.1

- (d) (i) Calculate the enthalpy of vaporization for gallium given the above data.

This is just products—reactants, so $277.1 - 5.60 = 271.5 \text{ kJ mol}^{-1}$

- (ii) The enthalpy of vaporization for gallium is greater than its enthalpy of fusion. Explain why in terms of intermolecular forces.

Enthalpy of vaporization describes the amount of heat necessary to completely break any attractions between gallium atoms, as ideal gas particles have no attractions between each other. That amount of energy will be greater than the amount necessary to simply weaken

them, which is what occurs in the solid to liquid phase change (quantified by the enthalpy of fusion).

- (e) Given your answer to (d)(i) and that $\Delta S^\circ = 113.4 \text{ J mol}^{-1} \text{ K}^{-1}$ for the boiling of gallium, what is the boiling point of the gallium?

During any phase change, $\Delta G^\circ = 0$. With that information, and converting the units to match:

$$\Delta G^\circ = \Delta H^\circ - T\Delta S^\circ$$

$$0 = 271.5 \text{ kJ mol}^{-1} - T(0.1134 \text{ kJ mol}^{-1} \text{ K}^{-1})$$

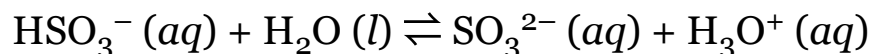
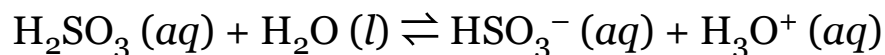
$$T = 2394 \text{ K}$$

- (f) After the gallium is fully converted to a gas, it continues to be heated. What would you expect to be true about the velocity distribution of the gaseous gallium atoms as the temperature increases?

As the gas is heated, the average energy in the gas will increase, causing a greater distribution of potential velocities for the gaseous atoms.

4. Sulfurous acid, H_2SO_3 , is a weak diprotic acid.

- (a) Write out the acid-base reactions which occur when H_2SO_3 is mixed with water.



- (b) A 0.50 M solution of H_2SO_3 has a pH of 1.10. Determine K_{a1} for H_2SO_3 .

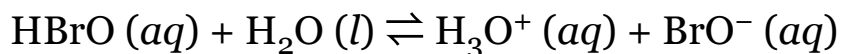
For a diprotic acid, only the first dissociation contributes significantly to the pH of the resulting solution. In this case, if $\text{pH} = 1.10$, then $-\log[\text{H}_3\text{O}^+] = 1.10$ and $[\text{H}_3\text{O}^+] = 0.079 \text{ M}$. Keeping in mind that $[\text{H}_3\text{O}^+] = [\text{HSO}_3^-]$, we get:

$$K_{\text{a}1} = \frac{[\text{H}_3\text{O}^+][\text{HSO}_3^-]}{[\text{H}_2\text{SO}_3]} \qquad K_{\text{a}1} = \frac{(0.079)^2}{(0.50 - 0.079)} \qquad K_{\text{a}1} = 1.5 \times 10^{-2}$$

Note that if the -0.079 is ignored in the denominator, it leads to the incorrect answer. The fact that some of the H_2SO_3 dissociates must be considered!

5. Hypobromous acid, HBrO , is a weak monoprotic acid with a K_{a} value of 2.0×10^{-9} at 25°C .

- (a) Write out the equilibrium reaction of hypobromous acid with water, identifying any conjugate acid/base pairs present.



Conjugate pairs: HBrO and BrO^- , H_2O and H_3O^+

- (b) (i) What would be the percent dissociation of a 0.50 M solution of hypobromous acid?

Percent dissociation describes what percentage of the acid ions will dissociate—that is, give up their hydrogen ions to form the conjugate base of BrO^- . First, you need the K_{a} expression:

$$K_{\text{a}} = \frac{[\text{H}_3\text{O}^+][\text{BrO}^-]}{[\text{HBrO}]}$$

For every proton that transfers, one BrO^- ion and one H_3O^+ ion are created, meaning those numbers will be identical and can both be replaced with x . The amount of dissociated ions will be insignificant compared to the amount of undissociated acid molecules in solution with such a small K_a , so you can ignore any change in the HBrO concentration.

$$2.0 \times 10^{-9} = \frac{x^2}{0.50 \text{ M}}$$

$$x = 3.2 \times 10^{-5} \text{ M}$$

Finally, to calculate percent dissociation, take the concentration of the conjugates created and divide it by the original concentration of the acid.

$$\% \text{ Dissociation} = \frac{3.2 \times 10^{-5} \text{ M}}{0.50 \text{ M}} \times 100\% = 6.4 \times 10^{-3} \%$$

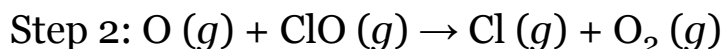
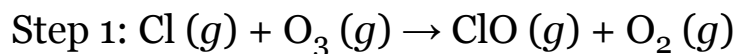
(ii) If the 0.50 M solution were diluted, what would happen to the percent dissociation of the HBrO ? Why?

The forward reaction, HBrO donating a proton to water, can occur easily regardless of the acid's concentration. This is because the vast majority of the HBrO remains undissociated, and there are always a very large amount of water molecules available to accept a proton from the acid.

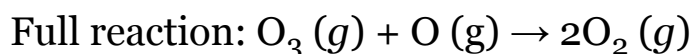
However, as the acid becomes less concentrated via dilution, less and less of the conjugates (BrO^- and H_3O^+) are created, making the reverse reaction increasingly less likely to happen, especially as the conjugates are surrounded by HBrO and water molecules. This means more of the conjugates will stay dissociated (they have a hard time “finding” each other to make the reaction

occur), which increases the percent dissociation of the acid.

6. Chlorofluorocarbons are byproducts of many different processes that are known to be dangerous to the environment as both as a greenhouse gas, as well as an agent for ozone (O_3) depletion. The accepted mechanism for the latter is:



- (a) Write out the full reaction with the above elementary steps, and identify all catalysts and intermediates.

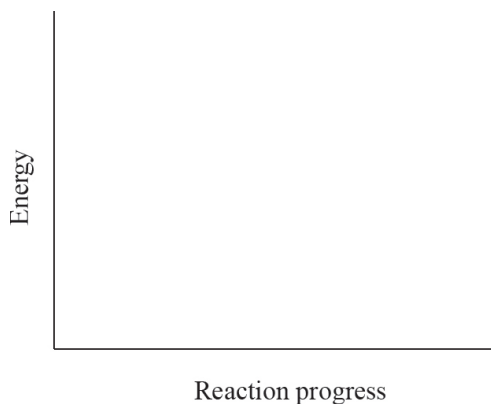


Catalyst: $\text{Cl} (g)$ Intermediate: $\text{ClO} (g)$

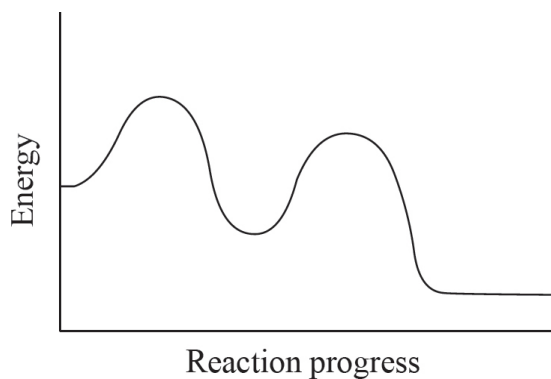
- (b) Describe two ways by which a catalyst can reduce the activation energy of a reaction.

A catalyst can either stabilize an existing intermediate, or it can cause the creation of a new intermediate. Both of these can lower the overall activation energy for the reaction, speeding it up.

- (c) Both elementary steps in the reaction above are exothermic. On the axes provided, draw a potential reaction mechanism which supports this.

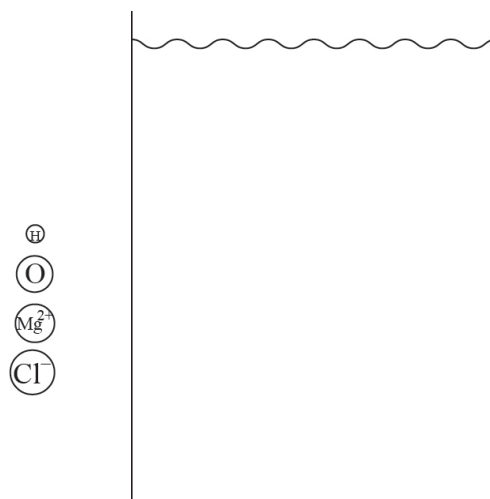


The most important thing in the diagram below is that the energy level decreases after each step.

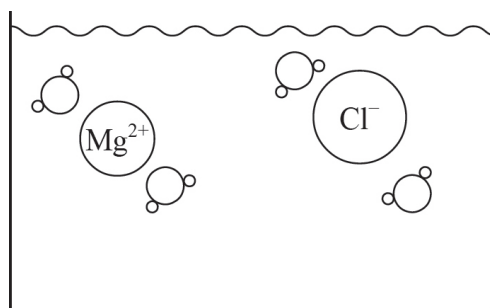


7. A stock solution of 2.0 M MgCl_2 is dissolved in water.

- (a) (i) In the beaker below, draw a particulate diagram that represents MgCl_2 dissolved in water. The approximate sizes of each atom/ion are provided for you. Your diagram should include at least four water molecules, which should be correctly oriented compared to the ions dissolved in solution.



Note that the negative (oxygen) dipoles on the water are attracted to the positive magnesium cations, while the positive (hydrogen) dipoles on the water attract to the negative chloride anions.



- (ii) Why are the chloride ions from (a)(i) larger than the magnesium ions?

The chlorine atom gains a single electron to become the chloride ion, filling its third energy level. The magnesium atom loses two electrons to form the magnesium ion, leaving it with only two full energy levels.

- (b) (i) A student wishes to make up 500 mL of 0.50 M $MgCl_2$ for an experiment. Explain the best method of doing so utilizing a graduated cylinder and a volumetric flask. Assume $MgCl_2$ is fully soluble.

For dilutions, use $M_1V_1 = M_2V_2$ to figure out the necessary volume of stock solution.

$$(2.0\text{ }M)(V_1) = (0.50\text{ }M)(500.\text{ mL})$$

$$V_1 = 125\text{ mL}$$

Using a graduated cylinder, add 125 mL of 2.0 *M* MgCl₂ to the volumetric flask. Next, dilute to the mark in the neck of the flask with distilled water before capping and inverting the flask several times to ensure a fully mixed solution.

- (ii) What are the concentrations of the Mg²⁺ and Cl⁻ ions in the new solution?

The equation for the dissociation is: $\text{MgCl}_2\text{ (s)} \rightarrow \text{Mg}^{2+}\text{ (aq)} + 2\text{Cl}^{-}\text{ (aq)}$

Thus, the magnesium ion concentration will be the same as the overall solution, and the chloride ion concentration will be twice that: $[\text{Mg}^{2+}] = 0.50\text{ }M$, $[\text{Cl}^{-}] = 1.0\text{ }M$

Part III

About the AP Chemistry Exam

THE STRUCTURE OF THE AP CHEMISTRY EXAM

The AP Chemistry Exam is a three-hour-long, two-section test that attempts to cover the material you would learn in a college first-year chemistry course. The first section is a 90-minute, 60-question multiple-choice section. As of 2015, the second section is 105 minutes and consists of three long-form free-response questions and four short-form free-response questions.

The multiple-choice section is scored by a computer, and the free-response questions are scored by a committee of high school and college teachers. The free-response questions are graded according to a standard set at the beginning of the grading period by the chief faculty consultants. Inevitably, the grading of Section II is never as consistent or accurate as the grading of Section I.

Take a look at the following data on how students scored on the May 2016 test:

Score (Meaning)	Percentage of Test Takers	Equivalent Grade in a first-year college course
5 (extremely qualified)	9.2%	A
4 (well qualified)	16.1%	A–, B+, B
3 (qualified)	28.1%	B–, C+, C
2 (possibly qualified)	24.9%	C–
1 (no recommendation)	21.7%	D

OVERVIEW OF CONTENT TOPICS

The concepts of the AP Chemistry Exam are broken down into six major themes defined by the College Board as the Big Ideas. Rather than learning multiple disparate topics (as has been done in the past), these Big Ideas interconnect principles within these topics that describe fundamental chemical phenomena. The six Big Ideas are as follows:

- Big Idea #1: The chemical elements are fundamental building materials of matter, and all matter can be understood in terms of arrangements of atoms. These atoms retain their identity in chemical reactions.
- Big Idea #2: Chemical and physical properties of materials can be explained by the structure and the arrangement of atoms, ions, or molecules and the forces between them.
- Big Idea #3: Changes in matter involve the rearrangement and/or reorganization of atoms and/or the transfer of electrons.
- Big Idea #4: Rates of chemical reactions are determined by details of the molecular collisions.
- Big Idea #5: The laws of thermodynamics describe the essential role of energy and explain and predict the direction of changes in matter.
- Big Idea #6: Any bond or intermolecular attraction that can be formed can be broken. These two processes are in a dynamic competition, sensitive to initial conditions and external perturbations.

This book has been arranged to teach chemistry topics grouped to each of these Big Ideas.

HOW AP EXAMS ARE USED

Different colleges use AP Exam scores in different ways, so it is important that you go to a particular college's website to determine how it uses AP Exam scores. The three items below represent the main ways in which AP Exam scores can be used.

- **College Credit.** Some colleges will give you college credit if you score well on an AP Exam. These credits count toward your graduation requirements, meaning that you can take fewer courses while in college. Given the cost of college, this could be quite a benefit, indeed.
- **Satisfy Requirements.** Some colleges will allow you to “place out” of certain requirements if you do well on an AP Exam, even if they do not give you actual college credits. For example, you might not need to take an introductory-level course, or perhaps you might not need to take a class in a certain discipline at all.
- **Admissions Plus.** Even if your AP Exam will not result in college credit or even allow you to place out of certain courses, most colleges will respect your decision to push yourself by taking an AP course or even an AP Exam outside of a course. A high score on an AP Exam shows mastery of more difficult content than is taught in many high school courses, and colleges may take that into account during the admissions process.

OTHER RESOURCES

There are many resources available to help you improve your score on the AP Chemistry Exam, not the least of which are your teachers. If you are taking an AP class, you may be able to get extra attention from your teacher, such as obtaining feedback on your free-response questions. If you are not in an AP course, reach out to a teacher who teaches chemistry, and ask if the teacher will review your free-response questions or otherwise help you with content.



Want to know which colleges are best for you? Check out The Princeton Review's College Advisor app to build your ideal college list and find your perfect college fit! Available in the iOS App Store and Google Play Store.



Another wonderful resource is **AP Students**, the official site of the AP Exams. The scope of the information at this site is quite broad and includes:

- Course Description, which includes details on what content is covered and sample multiple-choice and free-response questions
- Exam practice tips
- Free-response question prompts and scoring guidelines from previous AP Chemistry exams

The AP Students home page address is:
<https://apstudent.collegeboard.org>.

Finally, The Princeton Review offers tutoring and small group instruction. Our expert instructors can help you refine your strategic approach and add to your content knowledge. For more information, call 1-800-2REVIEW.

DESIGNING YOUR STUDY PLAN

As part of the Introduction, you identified some areas of potential improvement. Let's now delve further into your performance on Test

1, with the goal of developing a study plan appropriate to your needs and time commitment.

Read the answers and explanations associated with the multiple-choice questions (starting at [this page](#)). After you have done so, respond to the following:

- Review the Overview of Content Topics on [this page](#). Next to each topic, indicate your rank of the topic as follows: “1” means “I need a lot of work on this,” “2” means “I need to beef up my knowledge,” and “3” means “I know this topic well.”
- How many days/weeks/months away is your exam?
- What time of day is your best, most focused study time?
- How much time per day/week/month will you devote to preparing for your exam?
- When will you do this preparation? (Be as specific as possible: Mondays and Wednesdays from 3:00 P.M.–4:00 P.M., for example)
- Based on the answers above, will you focus on strategy (Part IV) or content ([Part V](#)) or both?
- What are your overall goals in using this book?

Part IV

Test-Taking Strategies for the AP Chemistry Exam

Chapter 1

How to Approach Multiple-Choice Questions

THE BASICS

Section I of the test is composed of 60 multiple-choice questions for which you are allotted 90 minutes. This part is worth 50% of your total score.

For this section, you will be given a periodic table of the elements along with a sheet that lists common chemistry formulas, and you may NOT use a calculator. The College Board says that this is because the new scientific calculators not only program and graph but also store information—and they are afraid you'll use this function to cheat!

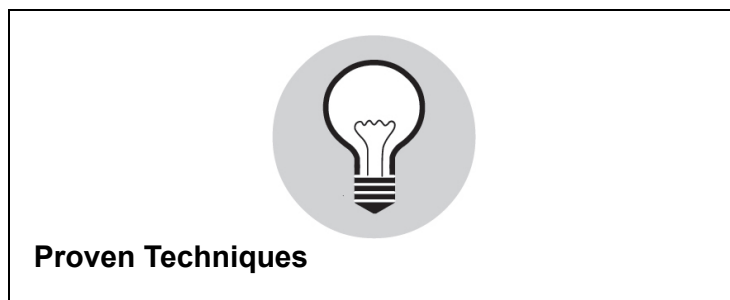
On the multiple-choice section, you receive 1 point for a correct answer. There is no penalty for leaving a question blank or getting a question wrong.

PACING

According to the College Board, the multiple-choice section of the AP Chemistry Exam covers more material than any individual student is expected to know. Nobody is expected to get a perfect or even near perfect score. What does that mean to you?

Use the Two-Pass System

Go through the multiple-choice section twice. The first time, do all the questions that you can get answers to immediately. That is, do the questions with little or no math and questions on chemistry topics in which you are well versed. Skip questions on topics that make you uncomfortable. Also, you want to skip the ones that look like number crunchers (even without a calculator, you may still be expected to crunch a few numbers). Circle the questions that you skip in your test booklet so you can find them easily during the second pass. Once you've done all the questions that come easily to you, go back and pick out the tough ones that you have the best shot at.



In general, the questions near the end of the section are tougher than the questions near the beginning. You should keep that in mind, but be aware that each person's experience will be different. If you can do acid-base questions in your sleep, but you'd rather have your teeth drilled than draw a Lewis diagram, you may find questions near the end of the section easier than questions near the beginning.

That's why the Two-Pass System is so handy. By using it, you make sure you get to see all the questions you can get right, instead of running out of time because you got bogged down on questions you couldn't do earlier in the test.

Don't Turn a Question into a Crusade!

Most people don't run out of time on standardized tests because they work too slowly. Instead, they run out of time because they spend half the test wrestling with two or three particular questions.

You should never spend more than a minute or two on any question. If a question doesn't involve calculation, then either you know the answer, you can make an educated guess, or you don't know the answer. Figure out where you stand on a question, make a decision, and move on.

Any question that requires more than two minutes worth of calculations probably isn't worth doing. Remember, skipping a question early in the section is a good thing if it means that you'll have time to get two correct answers later on.

GUESSING

You get one point for every correct answer on the multiple-choice section. Guessing randomly neither helps you nor hurts you. Educated guessing, however, will help you.

Use Process of Elimination (POE) to Find Wrong Answers

There is a fundamental weakness to a multiple-choice test. The test makers must show you the right answer, along with three wrong answers. Sometimes seeing the right answer is all you need. Other times you may not know the right answer, but you may be able to identify one or two of the answers that are clearly wrong. Here is where you should use Process of Elimination (POE) to take an educated guess.



Look at this hypothetical question.

1. Which of the following compounds will produce a purple solution when added to water?
 - (A) Brobogdium rabelide
 - (B) Diblythium perjuvenide
 - (C) Sodium chloride
 - (D) Carbon dioxide

You should have no idea what the correct answer is because two of these compounds are made up, but you do know something about the obviously wrong answers. You know that sodium chloride, (C), and carbon dioxide, (D), do not turn water purple. So, using POE, you have a 50 percent chance at guessing the correct answer. Now the odds are in your favor. Now you should guess.

Guess and Move On

Remember that you're guessing. Pondering the possible differences between brobogdium rabelide and diblythium perjuvenide is a waste of time. Once you've taken POE as far as it will go, pick your favorite letter and move on.

The multiple-choice section is the exact opposite of the free-response section. It's scored by a machine. There's no partial credit. The computer doesn't know, or care if you know, why an answer is correct. All the computer cares about is whether you blackened in the correct oval on your score sheet. You get the same number of points for picking (B) because you know that (A) is wrong and that (B) is a nicer letter than (C) or (D) as you would get for picking (B) because you fully understood the subtleties of an electrochemical process.

ABOUT CALCULATORS

You will NOT be allowed to use a calculator on this section. That shouldn't worry you. All it means is that there won't be any questions in the section that you'll need a calculator to solve.

Most of the calculation problems will have fairly user-friendly numbers—that is, numbers with only a couple of significant digits, or things like “11.2 liters of gas at STP” or “160 grams of oxygen” or “a temperature increase from 27°C to 127°C.” Sometimes these user-friendly numbers will actually point you toward the proper steps to take in your calculations.

Don't be afraid to make rough estimates as you do your calculations. Sometimes knowing that an answer is closer to 50 than to 500 will enable you to pick the correct answer on a multiple-choice test (if the answer choices are far enough apart). Once again, the rule against calculators works in your favor because the College Board will not expect you to do very precise calculations by hand.

REFLECT

Respond to the following questions:

- How long will you work on multiple-choice questions?
- How will you change your approach to multiple-choice questions?
- What is your multiple-choice guessing strategy?

Chapter 2

How to Approach Free-Response Questions

OVERVIEW OF THE FREE-RESPONSE SECTION

Section II is composed of seven free-response questions. You will be given 105 minutes to complete this section, which is worth 50 percent of your total score.

The first three free-response questions are longer, and are worth 10 points each. The College Board recommends you take about 22 minutes per question on these. The last four questions are much shorter, and only worth 4 points each. For these, the College Board recommends you take about 9 minutes each.

The suggested times are just that—suggestions. Your results may vary based on how comfortable you are with the topic being tested within each question. They do give you a good guideline, though. You probably shouldn't be spending more than 20 minutes on any of the first three questions. Don't be afraid to cut yourself off and come back to questions later if time allows.

CRACKING THE MATH PROBLEMS

You want to show the graders that you understand the math behind these chemistry problems, so here are some suggestions.

Show Every Step of Your Calculations on Paper

This section is the opposite of multiple-choice. You don't just get full credit for writing the correct answer. You get most of your points on this section for showing the process that got you to the answer. The graders give you partial credit when you show them that you know what you're doing. So even if you can do a calculation in your head, you should set it up and show it on the page.

By showing every step, or explaining what you're doing in words, you ensure that you'll get all the partial credit possible, even if you screw

up a calculation.

Include Units in All Your Calculations

Scientists like units in calculations. Units make scientists feel secure. You'll get points for including them and you may lose points for leaving them out.

Remember Significant Figures

You can lose 1 point per question if your answer is off by more than one significant figure. Without getting too bent out of shape about it, try to remember that a calculation is only as accurate as the least accurate number in it.

(For a detailed explanation of significant figures, see Chapter 9.)

The Graders Will Follow Your Reasoning, Even if You've Made a Mistake

Often, you are asked to use the result of a previous part of a problem in a later part. If you got the wrong answer in part (a) and used it in part (c), you can still get full credit for part (c), as long as your work is correct based on the number that you used. That's important, because it means that botching the first part of a question doesn't necessarily sink the whole question.

Remember the Mean!

So let's say that you could complete only parts (a) and (b) on the required equilibrium problem. That's 4 or 5 points out of 10, tops. Are you doomed? Of course not. You're above average. If this test is hard on you, it's probably just as hard on everybody else. Remember, you don't need anywhere near a perfect score to get a 5, and you can leave half the test blank and still get a 3!

STRATEGIES FOR CRACKING THE FREE-RESPONSE SECTION

This section is here to test whether you can translate chemistry into English. Most of the questions can be answered in two or three simple sentences, or with a simple diagram or two. Here are some tips for answering the seven free-response questions on Part II.

Show That You Understand the Terms Used in the Question

If you are asked why sodium and potassium have differing first ionization energies, the first thing you should do is tell them what ionization energy is. That's probably worth the first point of partial credit. Then you should tell them how the differing structures of the atoms make for differing ionization energies. That leads to the next tip.

Take a Step-by-Step Approach

Grading these tests is hard work. Breaking a question into parts in this way makes it easier on the grader, who must match your response to a set of guidelines he or she has been given that describe how to assign partial and full credit.



Each grader scores each test based on these rough guidelines that are established at the beginning of the grading period. For instance, if a grader has 3 points for the question about ionization energies, the points might be distributed the following way:

- one point for understanding ionization energy
- one point for explaining the structural difference between sodium and potassium
- one point for showing how this difference affects the ionization energy

You can get all three points for this question if the grader thinks that all three concepts are addressed implicitly in your answer, but by taking a step-by-step approach, you improve your chances of

explicitly addressing the things that a grader has been instructed to look for. Once again, grading these tests is hard work; graders won't know for sure if you understand something unless you tell them.

Write Neatly

This simple concept cannot be stressed enough: Write neatly, even if that means working at half-speed. You can't get points for answers if the graders can't understand them. Of course, this applies to the rest of the free-response section as well.

Graders Will Follow Your Reasoning, Even If You Make A Mistake

Just like in the multiple-choice section, you might be asked to use the result of a previous part of a problem in a later part. If you decide (incorrectly) that an endothermic reaction in part (a) is exothermic, you can still get full credit in part (c) for your wrong answer about the reaction's favorability, as long as your answer in (c) is correct based on an exothermic reaction.

REFLECT

Respond to the following questions:

- How much time will you spend on the short free-response questions? What about the long free-response questions?
- What will you do before you begin writing your free-response answers?
- Will you seek further help, outside of this book (such as a teacher, tutor, or AP Students), on how to approach the questions that you will see on the AP Chemistry Exam?

Part V

Content Review for the AP Chemistry Exam

Chapter 3

Big Idea #1: Atoms, Elements, and the Building Blocks of Matter

The chemical elements are fundamental building materials of matter, and all matter can be understood in terms of arrangements of atoms. These atoms retain their identity in chemical reactions.

THE PERIODIC TABLE

The most important tool you will use on this test is the Periodic Table of the Elements.

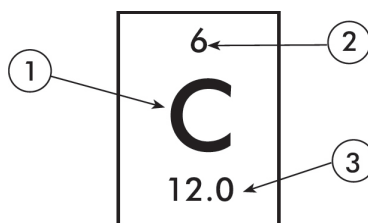
PERIODIC TABLE OF THE ELEMENTS																	
1 IA																	18 VIIIA
1 H 1.008	2 He 4.00																
3 Li 6.94	4 Be 9.01											5 B 10.81	6 C 12.01	7 N 14.01	8 O 16.00	9 F 19.00	10 Ne 20.18
11 Na 22.99	12 Mg 24.30	13 Al 26.98	14 Si 28.09	15 P 30.97	16 S 32.06	17 Cl 35.45	18 Ar 39.95										
19 K 39.10	20 Ca 40.08	21 Sc 44.96	22 Ti 47.90	23 V 50.94	24 Cr 52.00	25 Mn 54.94	26 Fe 55.85	27 Co 58.93	28 Ni 58.69	29 Cu 63.55	30 Zn 65.39	31 Ga 69.72	32 Ge 72.59	33 As 74.92	34 Se 78.96	35 Br 79.90	36 Kr 83.80
37 Rb 85.47	38 Sr 87.62	39 Y 88.91	40 Zr 91.22	41 Nb 92.91	42 Mo 95.94	43 Tc (98)	44 Ru 101.1	45 Rh 102.91	46 Pd 106.42	47 Ag 107.87	48 Cd 112.41	49 In 114.82	50 Sn 118.71	51 Sb 121.75	52 Te 127.60	53 I 126.91	54 Xe 131.29
55 Cs 132.91	56 Ba 137.33	57 *La 138.91	72 Hf 178.49	73 Ta 180.95	74 W 183.85	75 Re 186.21	76 Os 190.2	77 Ir 192.2	78 Pt 195.08	79 Au 196.97	80 Hg 200.59	81 Tl 204.38	82 Pb 207.2	83 Bi 208.98	84 Po (209)	85 At (210)	86 Rn (222)
87 Fr (223)	88 Ra 226.02	89 †Ac 227.03	104 Rf (261)	105 Db (262)	106 Sg (266)	107 Bh (264)	108 Hs (277)	109 Mt (268)	110 Ds (271)	111 Rg (272)							

*Lanthanide Series

58 Ce 140.12	59 Pr 140.91	60 Nd 144.24	61 Pm (145)	62 Sm 150.4	63 Eu 151.97	64 Gd 157.25	65 Tb 158.93	66 Dy 162.50	67 Ho 164.93	68 Er 167.26	69 Tm 169.93	70 Yb 173.04	71 Lu 174.97
90 Th 232.04	91 Pa 231.04	92 U 238.03	93 X (237)	94 Pu (244)	95 Am (243)	96 Cm (247)	97 Bk (247)	98 Cf (251)	99 Es (252)	100 Fm (257)	101 Md (258)	102 No (259)	103 Lr (262)

†Actinide Series

The Periodic Table gives you very basic but very important information about each element.



1. This is the **symbol** for the element; carbon, in this case. On the test, the symbol for an element is used interchangeably with the name of the element.

2. This is the **atomic number** of the element. The atomic number is the same as the number of protons in the nucleus of an element; it is also the same as the number of electrons surrounding the nucleus of an element when it is neutrally charged.
3. This number represents the average atomic mass of a single atom of carbon, measured in atomic mass units (amu). It also represents the average mass for a mole (see [this page](#)) of carbon atoms, measured in grams. Thus, one mole of carbon atoms has a mass of 12.01 g. This is called the **molar mass** of the element.

Bad Joke Alert

To prepare for the AP Chemistry Exam, familiarize yourself with the periodic table of elements by looking at it periodically.

The horizontal rows of the periodic table are called **periods**.

The vertical columns of the periodic table are called **groups**.



Groups can be numbered in two ways. The old system used Roman numerals to indicate groups. The new system simply numbers the groups from 1 to 18. While it is not important to know the specific group numbers, it is important to know the names of some groups.

Group IA/1 – Alkali Metals

Group IIA/2 – Alkaline Earth Metals

Group B/3-12 – Transition Metals

Group VIIA/17 – Halogens

Group VIIIA/18 – Noble Gases

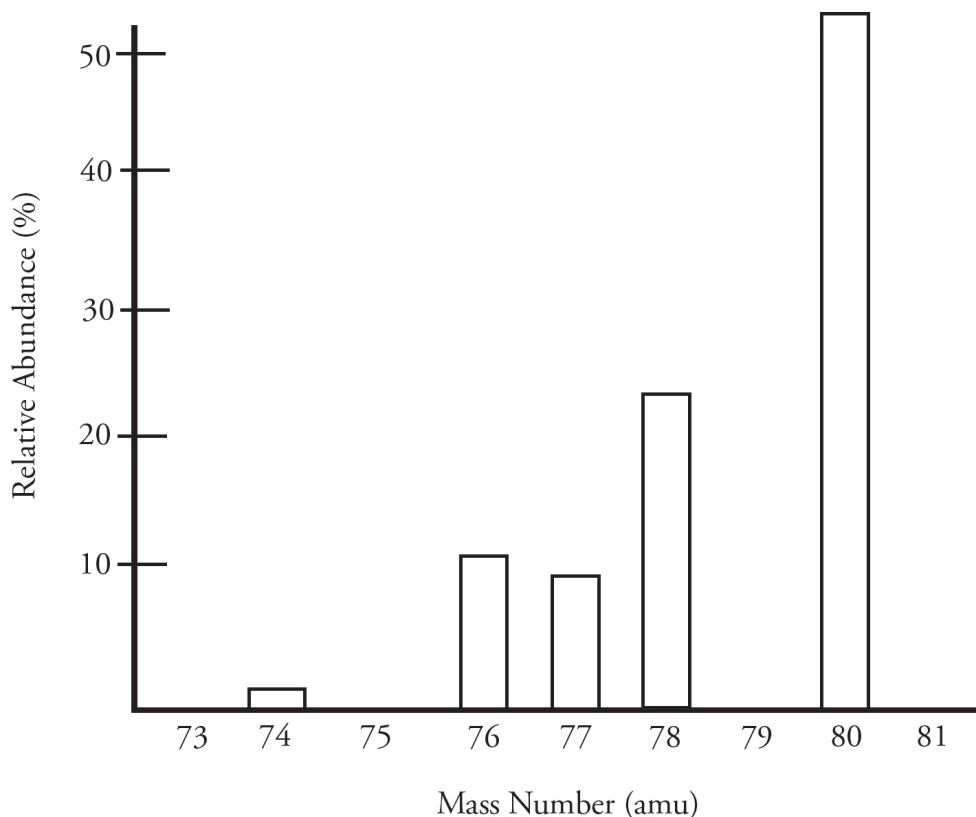
In addition, the two rows offset beneath the table are alternatively called the lanthanides and actinides, the rare Earth elements, or the

inner transition metals.

The identity of an atom is determined by the number of protons contained in its nucleus. The nucleus of an atom also contains neutrons. The mass number of an atom is the sum of its neutrons and protons. Electrons have significantly less mass than protons or neutrons and do not contribute to an element's mass.

Atoms of an element with different numbers of neutrons are called **isotopes**; for instance, carbon-12, which contains 6 protons and 6 neutrons, and carbon-14, which contains 6 protons and 8 neutrons, are isotopes of carbon. The molar mass given on the periodic table is the average of the mass numbers of all known isotopes weighted by their percent abundance.

The mass of various isotopes of an element can be determined by a technique called mass spectrometry. A mass spectrum of selenium looks like the following:



As you can see, the most abundant isotope of selenium has a mass of 80, but there are four other naturally occurring isotopes of selenium. The average atomic mass is the weighted average of all five isotopes of selenium shown on this spectra.

The molar mass of an element will give you a pretty good idea of the most common isotope of that element. For instance, the molar mass of carbon is 12.01 and about 99 percent of the carbon in existence is carbon-12.

MOLES

The mole is the most important concept in chemistry, serving as a bridge that connects all the different quantities that you'll come across in chemical calculations. The coefficients in chemical reactions tell you about the reactants and products in terms of moles, so most of the stoichiometry questions you'll see on the test will be exercises in converting between moles and grams, liters, molarities, and other units.

Moles and Molecules

Avogadro's number describes how many atoms are in a single mole of any given element. Much like a dozen is always 12, Avogadro's number is always 6.02×10^{23} . While it technically can be used to count anything, due to its extremely large value it is usually only used to count extraordinarily small things. Within the confines of this book, it will be used to count atoms, molecules, electrons, or ions, depending on the problem.

$$1 \text{ mole} = 6.022 \times 10^{23} \text{ particles}$$

$$\text{Moles} = \frac{\text{particles}}{(6.022 \times 10^{23})}$$

Moles and Grams

Moles and grams can be related using the atomic masses given in the periodic table. Atomic masses on the periodic table are given in terms of atomic mass units (amu); however, they also signify how many grams are present in one mole of an element. So, if 1 carbon atom has a mass of 12 amu, then 1 mole of carbon atoms has a mass of 12 grams.

You can use the relationship between amu and g/mol to convert between grams and moles by using the following equation:

$$\text{Moles} = \frac{\text{grams}}{\text{molar mass}}$$

Moles and Gases

We'll talk more about the ideal gas equation in Chapter 4, but for now, you should know that you can use it to calculate the number of moles of a gas if you know some of the gas's physical properties. All you need to remember at this point is that in the equation $PV = nRT$, n stands for moles of gas.

$$\text{Moles} = \frac{PV}{RT}$$

P = pressure (atm)

V = volume (L)

T = temperature (K)

R = the gas constant, 0.0821 L·atm/mol·K

The equation above gives the general rule for finding the number of moles of a gas. Many gas problems will take place at STP, or standard temperature and pressure, where $P = 1$ atmosphere and $T = 273$ K. At STP, the situation is much simpler and you can convert directly between the volume of a gas and the number of moles. That's because at STP, one mole of gas always occupies 22.4 liters.

$$\text{Moles} = \frac{\text{liters}}{(22.4 \text{ L/mol})}$$

Molarity

Molarity (M) expresses the concentration of a solution in terms of volume. It is the most widely used unit of concentration, turning up in calculations involving equilibrium, acids and bases, and electrochemistry, among others. When you see a chemical symbol in brackets on the test, that means they are talking about molarity. For instance, “[Na⁺]” is the same as “the molar concentration (molarity) of sodium ions.”

$$\text{Molarity } (M) = \frac{\text{moles of solute}}{\text{liters of solution}}$$

Percent Composition

Percent composition is the percent by mass of each element that makes up a compound. It is calculated by dividing the mass of each element or component in a compound by the total molar mass for the substance.

Calculate the percent composition of each element in calcium nitrate, $\text{Ca}(\text{NO}_3)_2$.

To do this, you need to first separate each element and count how many atoms are present. Subscripts outside of parentheses apply to all atoms inside of those parentheses.

Calcium: 1

Nitrogen: 2

Oxygen: 6

Then, multiply the number of atoms by the atomic mass of each element.

$$\text{Ca: } 40.08 \times 1 = 40.08$$

$$\text{N: } 14.01 \times 2 = 28.02$$

$$\text{O: } 16.00 \times 6 = 96.00$$

Adding up the masses of the individual elements will give you the atomic mass of that compound. Divide each individual mass by the total molar mass to get your percent composition.

$$40.08 + 28.02 + 96.00 = 164.10 \text{ g/mol}$$

$$\text{Ca: } 40.08/164.10 \times 100\% = 24.42\%$$

$$\text{N: } 28.02/164.10 \times 100\% = 17.07\%$$

$$\text{O: } 96.00/164.10 \times 100\% = 58.50\%$$

You can check your work at the end by making sure your percents add up to 100% (taking rounding into consideration).

$$24.42\% + 17.07\% + 58.50\% = 99.99\%. \text{ Close enough!}$$

Empirical and Molecular Formulas

You will also need to know how to determine the empirical and molecular formulas of a compound given masses or mass percents of the components of that compound. Remember that the empirical formula represents the simplest ratio of one element to another in a

compound (e.g., CH₂O), while the molecular formula represents the actual formula for the substance (e.g., C₆H₁₂O₆).

Let's take a look at the following example:

A compound is found to contain 56.5% carbon, 7.11% hydrogen, and 36.4% phosphorus.

a) Determine the empirical formula for the compound.

We start by assuming a 100 gram sample; this allows us to convert those percentages to grams. After we have that done, each element needs to be converted to moles.

$$\text{C: } 56.5\text{ g C} \times \frac{1 \text{ mol C}}{12.01 \text{ g C}} = 4.71 \text{ mol C}$$

$$\text{H: } 7.11 \text{ g H} \times \frac{1 \text{ mol H}}{1.01 \text{ g H}} = 7.04 \text{ mol H}$$

$$\text{P: } 36.4 \text{ g P} \times \frac{1 \text{ mol P}}{30.97 \text{ g P}} = 1.18 \text{ mol P}$$

We then divide each mole value by the lowest of the values. In this example, that would be the phosphorus. It is acceptable to round your answers if they are close (within 0.1) to a whole number.

$$\text{C: } \frac{4.71 \text{ mol}}{1.18 \text{ mol}} = 4$$

$$\text{H: } \frac{7.04 \text{ mol}}{1.18 \text{ mol}} = 6$$

$$\text{P: } \frac{1.18 \text{ mol}}{1.18 \text{ mol}} = 1$$

Those values become subscripts, so the empirical formula for the compound is $\text{C}_4\text{H}_6\text{P}$.

b) If the compound has a molar mass of 170.14 g/mol, what is its molecular formula?

First, we determine the molar mass of the empirical formula.

$$(12.01 \times 4) + (1.01 \times 6) + 30.97 = 85.07 \text{ g/mol}$$

Then, we divide that mass into the molar mass.

$$\frac{170.14}{85.07} = 2$$

Finally, multiply all subscripts in the empirical formula by that value. So, the molecular formula is $\text{C}_8\text{H}_{12}\text{P}_2$.

Electron Configurations and the Periodic Table

The positively charged nucleus is always pulling at the negatively charged electrons around it, and the electrons have potential energy that increases with their distance from the nucleus. It works the same way that the gravitational potential energy of a brick on the

third floor of a building is greater than the gravitational potential energy of a brick nearer to ground level.

The energy of electrons, however, is **quantized**. That's important. It means that electrons can exist only at specific energy levels, separated by specific intervals. It's kind of like if the brick in the building could be placed only on the first, second, or third floor of the building, but not in-between.

COULOMB'S LAW

The amount of energy that an electron has depends on its distance from the nucleus of the atom. This can be calculated using Coulomb's Law:

$$E = \frac{k(+q)(-q)}{r^{(2)}}$$

E = energy

k = Coulomb's constant

$+q$ = magnitude of the positive charge (nucleus)

$-q$ = magnitude of the negative charge (electron)

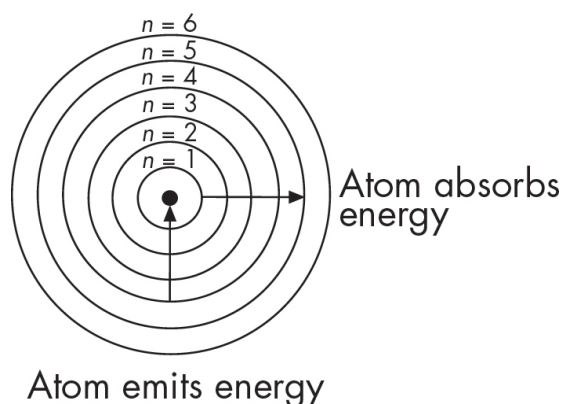
r = distance between the charges

While on the exam, you will not be required to mathematically calculate the amount of energy a given electron has, you should be able to qualitatively apply Coulomb's Law. Essentially, the greater the charge of the nucleus, the more energy an electron will have (as all electrons have the same amount of charge). Coulombic potential energy is considered to be 0 at a distance of infinity. The Coulombic potential energy for a 1s electron is lower (more negative) than that of say, a 3s electron. The amount of energy required to remove a 1s electron, thereby bringing its Coulombic potential energy to zero,

will thus be greater than the amount needed to remove a 3s electron. This removal energy is called the binding energy of the electron and is always a positive value.

The Bohr Model

Neils Bohr took the quantum theory and used it to predict that electrons orbit the nucleus at specific, fixed radii, like planets orbiting the Sun.



Each energy level is represented by a row on the periodic table. There are currently seven known energy levels, which correspond with $n = 1$ to $n = 7$. The closer an energy level is to an atom, the less energy electrons on that level have. While the Bohr model is not a perfect model of the atom, it serves as an excellent basis to understand atomic structure.

When atoms absorb energy in the form of **electromagnetic radiation**, electrons jump to higher energy levels. When electrons drop from higher to lower energy levels, atoms give off energy in the form of electromagnetic radiation.

The relationship between the change in energy level of an electron and the electromagnetic radiation absorbed or emitted is given below.

$$E = h\nu$$

E = energy change (in Joules)

h = Planck's constant (6.626×10^{-34} J s)

ν = frequency (in s^{-1})

The frequency and wavelength of electromagnetic radiation are inversely proportional. Combined with the energy and electromagnetic radiation equation, we can see that higher frequencies and shorter wavelengths lead to more energy.

Frequency and Wavelength

$$c = \lambda\nu$$

c = speed of light (2.998×10^8 ms^{-1})

ν = frequency (in s^{-1})

λ = wavelength (in m)

While frequency is always given in s^{-1} , wavelength is often given in nanometers, and needs to be converted to meters before being used in the above equation. An example is below.

$$350 \text{ nm} \times \frac{1 \times 10^{-9} \text{ m}}{1 \text{ nm}} = 3.50 \times 10^{-7} \text{ m}$$

Let's take a look at an example using the above equations.



How much energy does light with a wavelength of 695 nm have?

First, we have to convert wavelength to frequency. Remember, wavelength has to be in meters!

$$c = \lambda \nu$$

$$2.998 \times 10^8 \text{ m s}^{-1} = (6.95 \times 10^{-7} \text{ m})\nu$$

$$\nu = 4.31 \times 10^{14} \text{ s}^{-1}$$

Then, convert frequency to energy.

$$E = h\nu$$

$$E = (6.626 \times 10^{-34} \text{ J s})(4.31 \times 10^{14} \text{ s}^{-1})$$

$$E = 2.86 \times 10^{-19} \text{ J}$$

PHOTOELECTRON SPECTROSCOPY

If an atom is exposed to electromagnetic radiation at an energy level that exceeds the various binding energies of the electrons of that atom, the electrons can be ejected. The amount of energy necessary to do that is called the **ionization energy** for that electron. For the purposes of this exam, ionization energy and binding energy can be considered to be synonymous terms. When examining the spectra for electrons from a single atom or a small number of atoms, this energy is usually measured in electronvolts, eV ($1 \text{ eV} = 1.60 \times 10^{-19} \text{ Joules}$). If moles of atoms are studied, the unit for binding energy is usually either kJ/mol or MJ/mol.

All energy of the incoming radiation must be conserved and any of that energy that does not go into breaking the electron free from the

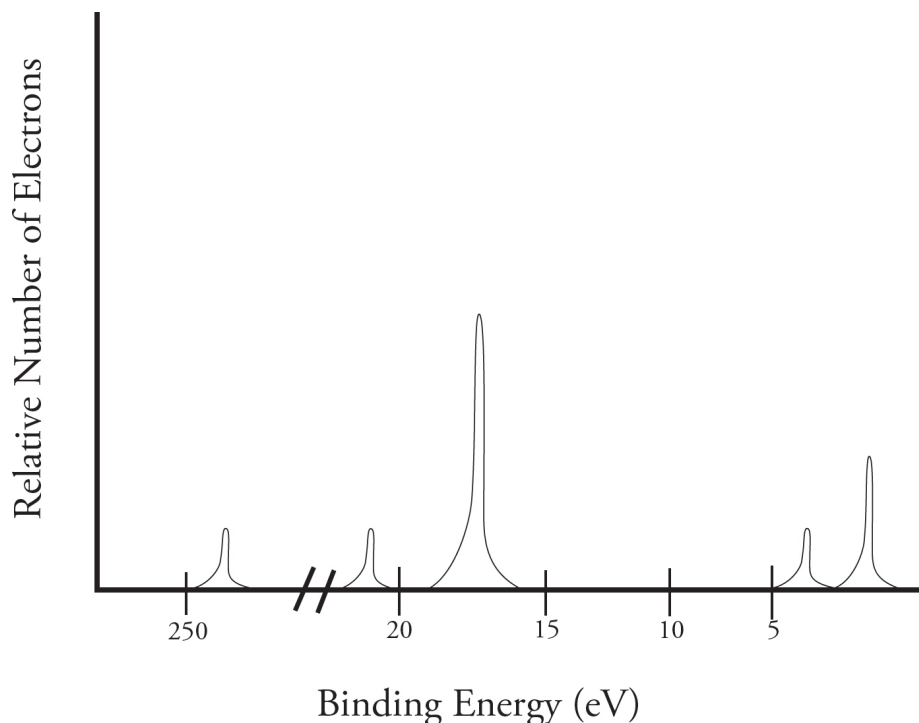
nucleus will be converted into **kinetic energy** (the energy of motion) for the ejected electron. So:

$$\text{Incoming Radiation Energy} = \text{Binding Energy} + \text{Kinetic Energy (of the ejected electron)}$$

The faster an ejected electron is going, the more kinetic energy it has. Electrons that were originally further away from the nucleus require less energy to eject, and thus will be moving faster. So, by examining the speed of the ejected electrons, we can determine how far they were from the nucleus of the atom in the first place. Usually, it takes electromagnetic radiation in either the visible or ultraviolet range to cause electron emission, while radiation in the infrared range is often used to study chemical bonds. Radiation in the microwave region is used to study the shape of molecules.

Spectra

If the amount of ionization energy for all electrons ejected from a nucleus is charted, you get what is called a **photoelectron spectra** (PES) that looks like the following:



The y -axis describes the relative number of electrons that are ejected from a given energy level, and the x -axis shows the binding energy of those electrons. Unlike most graphs, binding energy (ionization energy) decreases going from left to right in a PES. The spectra on the previous page is for sulfur.

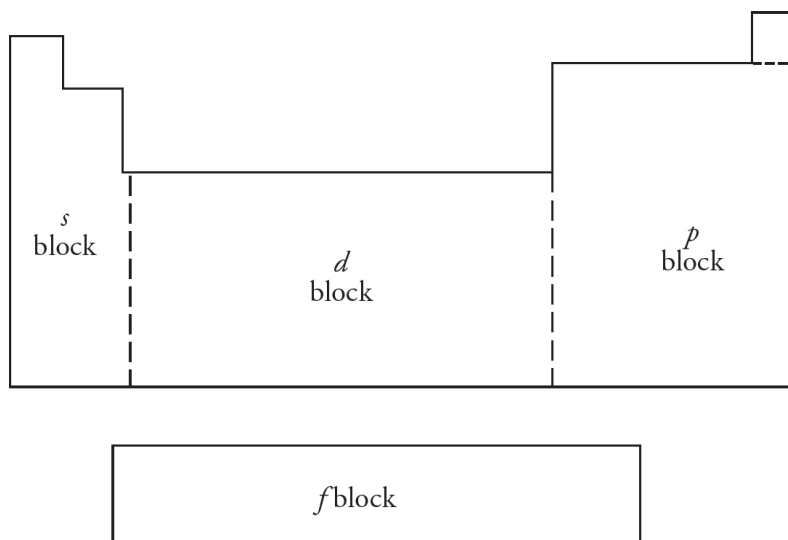
Each section of peaks in the PES represents a different energy level. The number of peaks in a section shows us that not all electrons at $n = 2+$ are located the same distance from the nucleus. With each energy level, there are **subshells**, which describe the shape of the space the electron can be found in (remember, we are in three-dimensional space here). The Bohr model is limited to two dimensions and does not represent the true positions of electrons due to that reason. Electrons do not orbit the nucleus as planets orbit the Sun. Instead, they are found moving about in a certain area of space (the subshell) a given distance (the energy level) away from the nucleus.

In all energy levels, the first subshell is called the s -subshell and can hold a maximum of two electrons. The second subshell is called the p -subshell and can hold a maximum of six electrons. In the spectra on the previous page, we can see the peak for the p -subshell in energy level 2 is three times higher than that of the s -subshell. The relative height of the peaks helps determine the number of electrons in that subshell.

In the area for the third energy level, the p -subshell peak is only twice as tall as the s -subshell. This indicates there are only four electrons in the p -subshell of this particular atom.

Electron Configuration

Studying the PES of elements allows scientists to understand more about the structure of the atom. In addition to the s and p subshells, two others exist: d (10 electrons max) and f (14 electrons max). The periodic table is designed so that each area is exactly the length of one particular subshell.



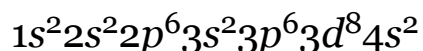
As you can see, the first two groups (plus helium) are in what is called the *s*-block. The groups on the right of the table are in the *p*-block, while transition metals make up the *d*-block. The inner transition metals below the table inhabit the *f*-block. The complete description of the energy level and subshell that each electron on an element inhabits is called its **electron configuration**.

Example 1: Determine the electron configuration for sulfur.

Sulfur has 16 electrons. The first two go into energy level 1 subshell *s*; this is represented by $1s^2$. The next two go into energy level 2 subshell *s*— $2s^2$. Six more fill energy level *p* ($2p^6$) then two more go into 3*s* ($3s^2$) and the final four enter into 3*p* ($3p^4$). So the final configuration is $1s^2 2s^2 2p^6 3s^2 3p^4$.

Using the periodic table as a reference can allow for the determination of any electron configuration. One thing to watch out for is that when entering the *d*-block (transition metals), the energy level drops by 1. The why behind that isn't important right now, but you should still be able to apply that rule.

Example 2: Determine the electron configuration for nickel.



Electron configurations can also be written “shorthand” by replacing parts of them with the symbol for the noble gas at the end of the highest energy level which has been filled.

For example, the shorthand notation for Si is $[\text{Ne}]3s^23p^2$ and the shorthand for nickel would be $[\text{Ar}]4s^23d^8$.

CONFIGURATION RULES

The Aufbau Principle

The **Aufbau principle** states that when building up the electron configuration of an atom, electrons are placed in orbitals, subshells, and shells in order of increasing energy.

The Pauli Exclusion Principle

The **Pauli Exclusion Principle** states that the two electrons which share an orbital cannot have the same spin. One electron must spin clockwise, and the other must spin counterclockwise.

Hund's Rule

Hund's Rule says that when an electron is added to a subshell, it will always occupy an empty orbital if one is available. Electrons always occupy orbitals singly if possible and pair up only if no empty orbitals are available.

Watch how the $2p$ subshell fills as we go from boron to neon.

	1s	2s	2p		
Boron	↑↓	↑↓	↑		
Carbon	↑↓	↑↓	↑	↑	
Nitrogen	↑↓	↑↓	↑	↑	↑
Oxygen	↑↓	↑↓	↑↓	↑	↑
Fluorine	↑↓	↑↓	↑↓	↑↓	↑
Neon	↑↓	↑↓	↑↓	↑↓	↑↓

PREDICTING IONIC CHARGES

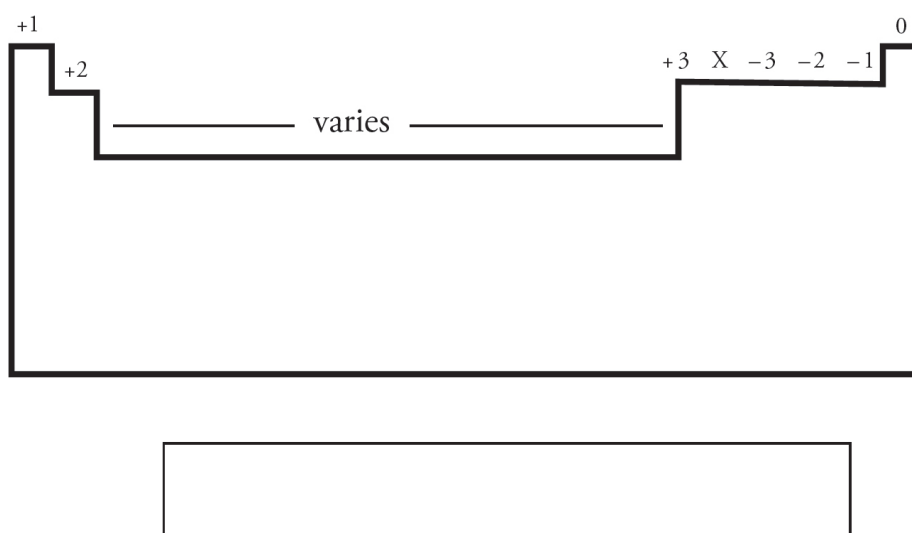
One of the great rules of chemistry is that the most stable configurations from an energy standpoint are those in which the outermost energy level is full. For anything in the *s* or *p* blocks, that means achieving a state in which there are eight electrons in the outermost shell (2 in the *s* subshell and 6 in the *p* subshell). The electrons in the outermost *s* and *p* subshells are called the **valence electrons** of that atom.

It's tempting to say that fluorine atoms "want" one electron or are "happy" with a full valence shell, but in reality, atoms don't have feelings and don't want anything! Instead, say fluorine atoms need only attract one electron to have a stable energy level. Assigning feelings and desires to inanimate particles won't do you any favors on the exam!

Elements that are close to a full energy level, such as the halogens or those in the oxygen group, tend to gain electrons to achieve a stable configuration. An **ion** is an atom which has either gained or lost electrons, while the number of protons and neutrons remains constant. Halogens need only gain one electron to achieve a stable configuration, and as such, typically form ions with a charge of negative one. F^- , Cl^- , etc. Any particle with more electrons than protons is called an **anion** (negatively charged ion). Those elements

in the oxygen group need two electrons for stability; thus, they have a charge of negative 2.

On the other end of the table, the alkali metals can most easily achieve a full valence shell by losing a single electron, rather than by gaining seven. So, they will form positively charged ions (**cations**), which have more protons than electrons. Alkali metals typically have a charge of +1, alkaline Earth metals of +2, and so forth. The table below gives a good overview of the common ionic charges for various groups:



In general, transition metals can form ions of varying charges. All transition metals lose electrons to form cations, but how many electrons they lose will vary depending on the compound they are in. For example, in CuBr_2 , we know what the charge on the bromide is negative one, and with two bromides the total negative charge present in the compound is -2 . The total compound charge must be zero for the compound to be stable, so it stands to reason that the charge on the single copper ion is +2. That compound would be called copper (II) bromide. However, in CuBr , the charge on the copper would only need to be +1 to balance the single bromide, so that compound would be called copper (I) bromide.

There are a few transition metals that only form ions with one possible charge. Two important examples are zinc, which always forms ions with a charge of +2, and silver, which always forms ions with a charge of +1.

There's also a slightly different rule as to how transition metals lose electrons when forming those cations. Transition metals, when losing electrons, will lose their higher-level *s* electrons before losing any of the lower-level *d* electrons. Thus, while an iron atom has a configuration of $[\text{Ar}]4s^23d^6$, an iron ion that has lost two electrons (Fe^{2+}) would have a configuration of $[\text{Ar}]3d^6$. If it were to lose further electrons, those would then come from the *d*-orbital.

Names and Theories

Dalton's Elements

In the early 1800s, John Dalton presented some basic ideas about atoms that we still use today. He was the first to say that there are many different kinds of atoms, which he called elements. He said that these elements combine to form compounds and that these compounds always contain the same ratios of elements. Water (H_2O), for instance, always has two hydrogen atoms for every oxygen atom. He also said that atoms are never created or destroyed in chemical reactions.

Development of the Periodic Table

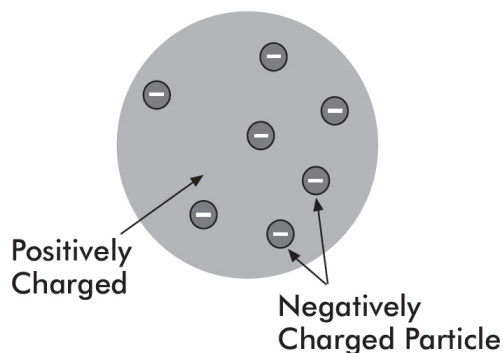
In 1869, Dmitri Mendeleev and Lothar Meyer independently proposed arranging the elements into early versions of the periodic table, based on the trends of the known elements.

Thomson's Experiment

In the late 1800s, J. J. Thomson watched the deflection of charges in a cathode ray tube and put forth the idea that atoms are composed of positive and negative charges. The negative charges were called electrons, and Thomson guessed that they were sprinkled throughout

the positively charged atom like chocolate chips sprinkled throughout a blob of cookie dough.

The Plum Pudding Model of an Atom

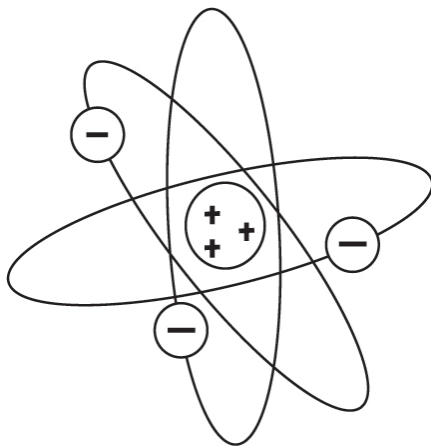


Millikan's Experiment

Robert Millikan was able to calculate the charge on an electron by examining the behavior of charged oil drops in an electric field.

Rutherford's Experiment

In the early 1900s, Ernest Rutherford fired alpha particles at gold foil and observed how they were scattered. This experiment led him to conclude that all of the positive charge in an atom was concentrated in the center and that an atom is mostly empty space. This led to the idea that an atom has a positively charged nucleus, which contains most of the atom's mass, and that the tiny, negatively charged electrons travel around this nucleus.



The Heisenberg Uncertainty Principle

Werner Heisenberg said that it is impossible to know both the position and momentum of an electron at a particular instant. In terms of atomic structure, this means that electron orbitals do not represent specific orbits like those of planets. Instead, an electron orbital is a probability function describing the possibility that an electron will be found in a region of space.

PERIODIC TRENDS

You can make predictions about certain behavior patterns of an atom and its electrons based on the position of the atom in the periodic table. All the periodic trends can be understood in terms of three basic rules.

1. Electrons are attracted to the protons in the nucleus of an atom.
 - a. The closer an electron is to the nucleus, the more strongly it is attracted.
 - b. The more protons in a nucleus, the more strongly an electron is attracted.
2. Electrons are repelled by other electrons in an atom. So, if other electrons are between a valence electron and the nucleus, the valence electron will be less attracted to the nucleus. That's called shielding.
3. Completed shells (and to a lesser extent, completed subshells) are very stable. Atoms will add or subtract valence electrons to create complete shells if possible.

The atoms in the left-hand side of the periodic table are called metals. Metals give up electrons when forming bonds. Most of the elements in the table are metals. The elements in the upper right-hand portion of the table are called nonmetals. Nonmetals generally gain electrons when forming bonds. The metallic character of the elements decreases as you move from left to right across the periodic table. The elements in the borderline between metal and nonmetal, such as silicon and arsenic, are called metalloids.

Atomic Radius

The atomic radius is the approximate distance from the nucleus of an atom to its valence electrons.

Moving from Left to Right Across a Period (Li to Ne, for Instance), Atomic Radius Decreases

Moving from left to right across a period, protons are added to the nucleus, so the valence electrons are more strongly attracted to the nucleus; this decreases the atomic radius. Electrons are also being added, but they are all in the same shell at about the same distance from the nucleus, so there is not much of a shielding effect.

Moving Down a Group (Li to Cs, for Instance), Atomic Radius Increases

Moving down a group, shells of electrons are added to the nucleus. Each shell shields the more distant shells from the nucleus and the valence electrons get farther away from the nucleus. Protons are also being added, but the shielding effect of the negatively charged electron shells cancels out the added positive charge.

Cations (Positively Charged Ions) Are Smaller than Atoms

Generally, when electrons are removed from an atom to form a cation, the outer shell is lost, making the cation smaller than the atom. Also, when electrons are removed, electron–electron repulsions are reduced, allowing all of the remaining valence electrons to move closer to the nucleus.

Anions (Negatively Charged Ions) Are Larger than Atoms

When an electron is added to an atom, forming an anion, electron–electron repulsions increase, causing the valence electrons to move farther apart, which increases the radius.

Ionization Energy

Electrons are attracted to the nucleus of an atom, so it takes energy to remove an electron. The energy required to remove an electron from an atom is called the first ionization energy. Once an electron has been removed, the atom becomes a positively charged ion. The energy required to remove the next electron from the ion is called the second ionization energy, and so on.

Moving from Left to Right Across a Period, Ionization Energy Increases

Moving from left to right across a period, protons are added to the nucleus, which increases its positive charge. For this reason, the negatively charged valence electrons are more strongly attracted to the nucleus, which increases the energy required to remove them.

Moving Down a Group, Ionization Energy Decreases

Moving down a group, shells of electrons are added to the nucleus. Each inner shell shields the more distant shells from the nucleus, reducing the pull of the nucleus on the valence electrons and making them easier to remove. Protons are also being added, but the shielding effect of the negatively charged electron shells cancels out the added positive charge.

The Second Ionization Energy Is Greater than the First Ionization Energy

When an electron has been removed from an atom, electron–electron repulsion decreases and the remaining valence electrons move closer to the nucleus. This increases the attractive force between the electrons and the nucleus, increasing the ionization energy.

As Electrons Are Removed, Ionization Energy Increases Gradually Until a Shell Is Empty, Then Makes a Big Jump

- For each element, when the valence shell is empty, the next electron must come from a shell that is much closer to the nucleus, making the ionization energy for that electron much larger than for the previous ones.
- For Na, the second ionization energy is much larger than the first.
- For Mg, the first and second ionization energies are comparable, but the third is much larger than the second.
- For Al, the first three ionization energies are comparable, but the fourth is much larger than the third.

You can also combine what you've learned about the math underlying energy to answer quantitative questions about ionization energy, such as the following:

Chlorine has a first ionization energy of 1251 kJ/mol. What wavelength of light would be necessary to ionize a single atom of chlorine?

The ionization energy is given in kJ/mol, and that needs to be converted to Joules before it can be used in the equations that relate energy and wavelength.

$$\frac{1251 \text{ kJ}}{\text{mol}} \times \frac{1 \text{ mol}}{6.02 \times 10^{23} \text{ atoms}} \times \frac{1000 \text{ J}}{1 \text{ kJ}} = 2.08 \times 10^{-18} \frac{\text{J}}{\text{atom}}$$

Then, energy can be converted to frequency.

$$E = h\nu$$
$$2.08 \times 10^{-18} \text{ J} = (6.626 \times 10^{-34} \text{ J s})\nu$$
$$\nu = 3.14 \times 10^{15} \text{ s}^{-1}$$

And finally, frequency can be converted to wavelength.

$$c = \lambda\nu$$
$$2.998 \times 10^8 \text{ m s}^{-1} = \lambda(3.14 \times 10^{15} \text{ s}^{-1})$$
$$\lambda = 9.55 \times 10^{-8} \text{ m}$$

Electronegativity

Electronegativity refers to how strongly the nucleus of an atom attracts the electrons of other atoms in a bond. Electronegativity is affected by two factors. The smaller an atom is, the more effectively its nuclear charge will be felt past its outermost energy level and the higher its electronegativity will be. Second, the closer an element is to having a full energy level, the more likely it is to attract the necessary electrons to complete that level. In general:

- Moving from left to right across a period, electronegativity increases.
- Moving down a group, electronegativity decreases.

The various periodic trends are summarized in the diagram below. The primary exception to these trends are the electronegativity values for the three smallest noble gases. As helium, neon, and argon do not form bonds, they have zero electronegativity. The larger noble gases, however, can form bonds under certain conditions and do follow the general trends as outlined in this section. This will be discussed in more detail in the next chapter.

- Atomic radius decreases
- Ionization energy increases
- Electronegativity increases

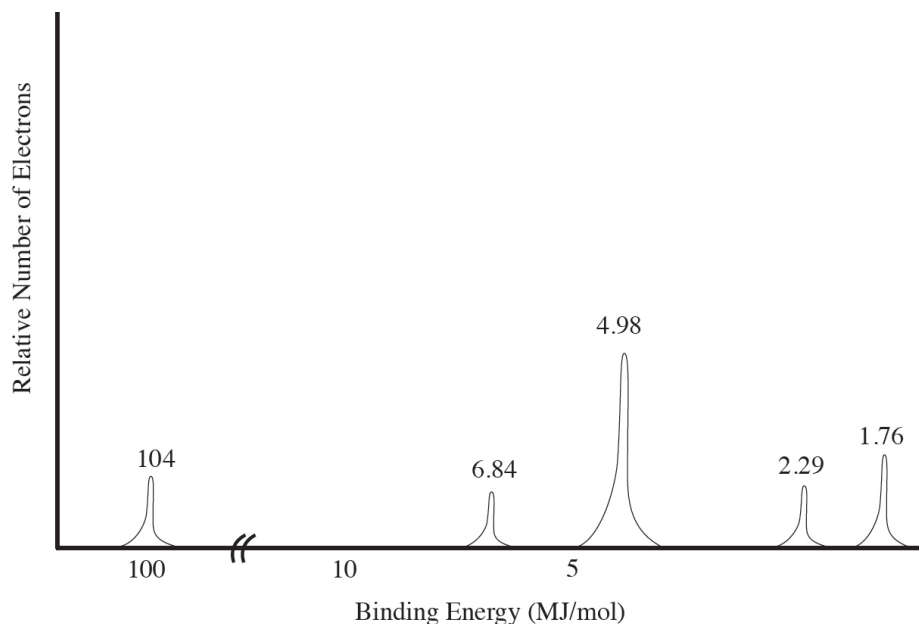
- Atomic radius increases
- Ionization energy decreases
- Electronegativity decreases

CHAPTER 3 QUESTIONS

[Click here](#) to download and print the PDF version of this exercise.

Multiple-Choice Questions

Use the PES spectra below to answer questions 1-4.



1. What element does this spectra represent?

- (A) Boron
- (B) Nitrogen
- (C) Aluminum
- (D) Phosphorus

2. Which peak represents the 2s subshell?

- (A) The peak at 104 MJ/mol
- (B) The peak at 6.84 MJ/mol

(C) The peak at 2.29 MJ/mol

(D) The peak at 1.76 MJ/mol

3. An electron from which peak would have the greatest velocity after ejection?

(A) The peak at 104 MJ/mol

(B) The peak at 6.84 MJ/mol

(C) The peak at 4.98 MJ/mol

(D) The peak at 1.76 MJ/mol

4. How many valence electrons does this atom have?

(A) 2

(B) 3

(C) 4

(D) 5

5. Why does an ion of phosphorus, P^{3-} , have a larger radius than a neutral atom of phosphorus?

(A) There is a greater Coulombic attraction between the nucleus and the electrons in P^{3-} .

(B) The core electrons in P^{3-} exert a weaker shielding force than those of a neutral atom.

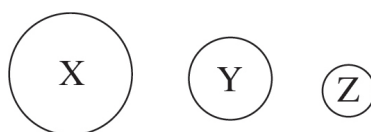
(C) The nuclear charge is weaker in P^{3-} than it is in P.

(D) The electrons in P^{3-} have a greater Coulombic repulsion than those in the neutral atom.

6. Which neutral atom of the following elements would have the most unpaired electrons?

- (A) Titanium
- (B) Manganese
- (C) Nickel
- (D) Zinc

7. The diagram below shows the relative atomic sizes of three different elements from the same period. Which of the following statements must be true?



- (A) The effective nuclear charge will be the greatest in element X.
 - (B) The first ionization energy will be greatest in element X.
 - (C) The electron shielding effect will be greatest in element Z.
 - (D) The electronegativity value will be greatest in element Z.
8. The first ionization energy for a neutral atom of chlorine is 1.25 MJ/mol and the first ionization energy for a neutral atom of argon is 1.52 MJ/mol. How would the first ionization energy value for a neutral atom of potassium compare to those values?
- (A) It would be greater than both because potassium carries a greater nuclear charge than either chlorine or argon.
 - (B) It would be greater than both because the size of a potassium atom is smaller than an atom of either chlorine or argon.
 - (C) It would be less than both because there are more electrons in potassium, meaning they repel each other more effectively and less energy is needed to remove one.
 - (D) It would be less than both because a valence electron of potassium is farther from the nucleus than one of either chlorine or argon.

9. Which of the following isoelectric species has the smallest radius?

(A) S^{2-}

(B) Cl^{-}

(C) Ar

(D) K^{+}

10. What is the most likely electron configuration for a sodium ion?

(A) $1s^2 2s^2 2p^5$

(B) $1s^2 2s^2 2p^6$

(C) $1s^2 2s^2 2p^6 3s^1$

(D) $1s^2 2s^2 2p^5 3s^2$

11. Which of the following statements is true regarding sodium and chlorine?

(A) Sodium has greater electronegativity and a larger first ionization energy.

(B) Sodium has a larger first ionization energy and a larger atomic radius.

(C) Chlorine has a larger atomic radius and a greater electronegativity.

(D) Chlorine has greater electronegativity and a larger first ionization energy.

12. An atom of silicon in its ground state is subjected to a frequency of light that is high enough to cause electron ejection. An electron from which subshell of silicon would have the highest kinetic energy after ejection?

- (A) $1s$
- (B) $2p$
- (C) $3p$
- (D) $4s$

13. The wavelength range for infrared radiation is 10^{-5} m, while that of ultraviolet radiation is 10^{-8} m. Which type of radiation has more energy, and why?

- (A) Ultraviolet has more energy because it has a higher frequency.
- (B) Ultraviolet has more energy because it has a longer wavelength.
- (C) Infrared has more energy because it has a lower frequency.
- (D) Infrared has more energy because it has a shorter wavelength.

14. Which of the following nuclei has 3 more neutrons than protons? (Remember: The number before the symbol indicates atomic mass.)

- (A) ^{11}B
- (B) ^{37}Cl
- (C) ^{24}Mg
- (D) ^{70}Ga

15. Examining data obtained from mass spectrometry supports which of the following?

- (A) The common oxidation states of elements
- (B) Atomic size trends within the periodic table

(C) Ionization energy trends within the periodic table

(D) The existence of isotopes

16. In general, do metals or nonmetals from the same period have higher ionization energies? Why?

(A) Metals have higher ionization energies because they usually have more protons than nonmetals.

(B) Nonmetals have higher ionization energies because they are larger than metals and harder to ionize.

(C) Metals have higher ionization energies because there is less electron shielding than there is in nonmetals.

(D) Nonmetals have higher ionization energies because they are closer to having filled a complete energy level.

17. The ionization energies for an element are listed in the table below.

First	Second	Third	Fourth	Fifth
8 eV	15 eV	80 eV	109 eV	141 eV

Based on the ionization energy table, the element is most likely to be

(A) Sodium

(B) Magnesium

(C) Aluminum

(D) Silicon

Use the following information to answer questions 18-20.

The outermost electron of an atom has a binding energy of 2.5 eV. The atom is exposed to light of a high enough frequency to

cause exactly one electron to be ejected. The ejected electron is found to have a KE of 2.0 eV.

18. How much energy did photons of the incoming light contain?
- (A) 0.50 eV
 - (B) 0.80 eV
 - (C) 4.5 eV
 - (D) 5.0 eV
19. If the wavelength of the light were to be shortened, how would that effect the kinetic energy of the ejected electron?
- (A) A shorter wavelength would increase the kinetic energy.
 - (B) A shorter wavelength would decrease the kinetic energy.
 - (C) A shorter wavelength would stop all electron emissions completely.
 - (D) A shorter wavelength would have no effect on the kinetic energy of the ejected electrons.
20. If the intensity of the light were to be decreased (that is, if the light is made dimmer), how would that affect the kinetic energy of the ejected electron?
- (A) The decreased intensity would increase the kinetic energy.
 - (B) The decreased intensity would decrease the kinetic energy.
 - (C) The decreased intensity would stop all electron emissions completely.
 - (D) The decreased intensity would have no effect.
21. Nitrogen's electronegativity value is between those of phosphorus and oxygen. Which of the following correctly describes the relationship between the three values?

- (A) The value for nitrogen is less than that of phosphorus because nitrogen is larger, but greater than that of oxygen because nitrogen has a greater effective nuclear charge.
- (B) The value for nitrogen is less than that of phosphorus because nitrogen has fewer protons, but greater than that of oxygen because nitrogen has fewer valence electrons.
- (C) The value for nitrogen is greater than that of phosphorus because nitrogen has fewer electrons, but less than that of oxygen because nitrogen is smaller.
- (D) The value for nitrogen is greater than that of phosphorus because nitrogen is smaller, but less than that of oxygen because nitrogen has a smaller effective nuclear charge.
22. Which of the following ions would have the most unpaired electrons?

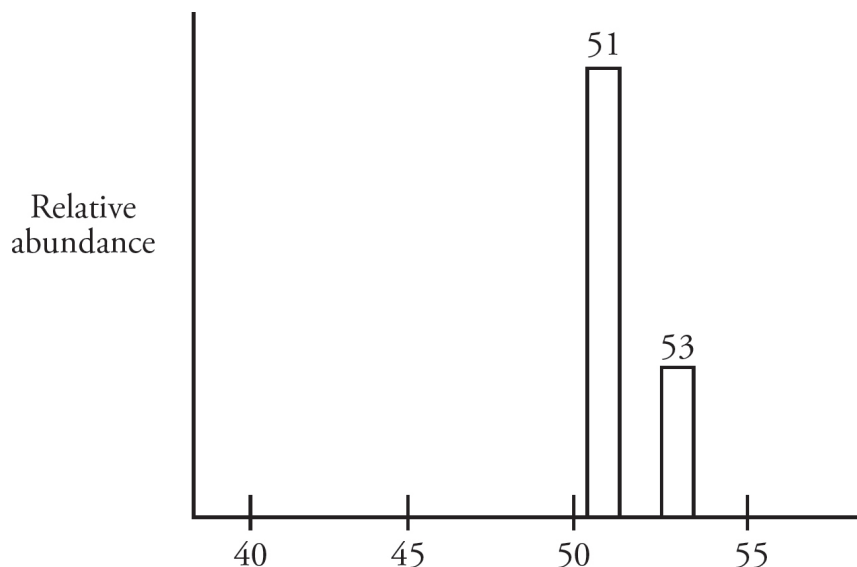
- (A) Mn^{2+}
- (B) Ni^{3+}
- (C) Ti^{2+}
- (D) Cr^{6+}

Free-Response Questions

1. Explain each of the following in terms of atomic and molecular structures and/or forces.
- (a) The first ionization energy for magnesium is greater than the first ionization energy for calcium.
- (b) The first and second ionization energies for calcium are comparable, but the third ionization energy is much greater.
- (c) There are three peaks of equal height in the PES of carbon, but on the PES of oxygen the last peak has a height twice as

high as all the others.

- (d) The first ionization energy for aluminum is lower than the first ionization energy for magnesium.

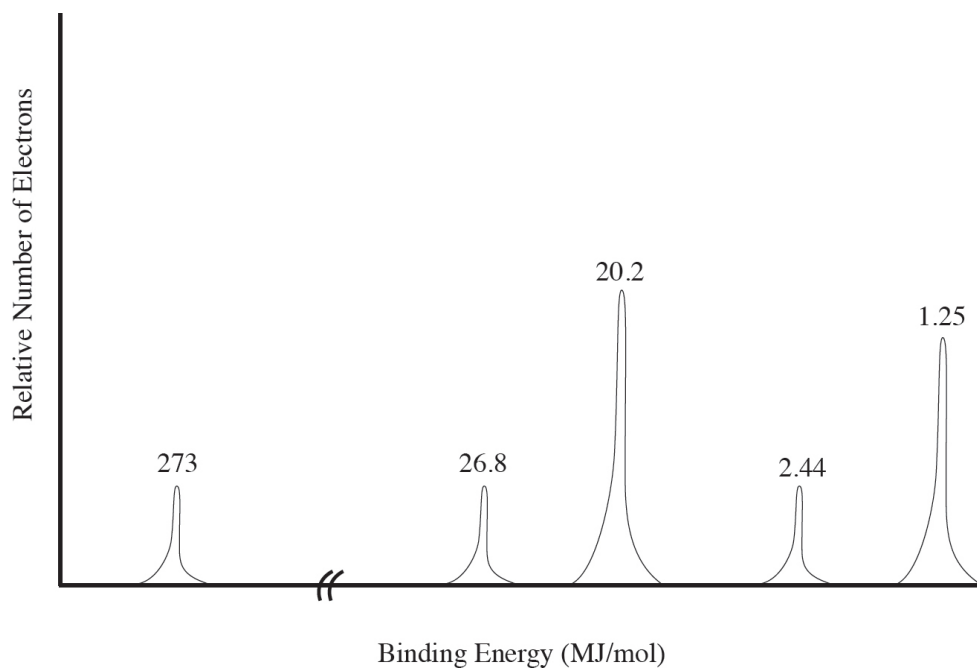


2. The above mass spectra is for the hypochlorite ion, ClO^- . Oxygen has only one stable isotope, which has a mass of 16 amu.
- (a) How many neutrons does the most common isotope of chlorine have?
 - (b) Using the spectra, calculate the average mass of a hypochlorite ion.
 - (c) Does the negative charge on the ion affect the spectra? Justify your answer.
 - (d) The negative charge in the ion is located around the oxygen atom. Speculate as to why.
3. The table below gives data on four different elements, in no particular order:

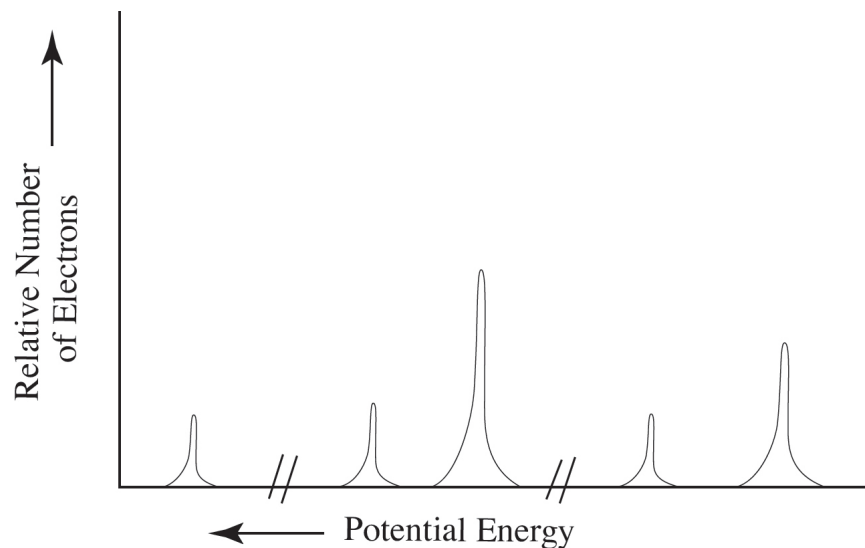
Carbon, Oxygen, Phosphorus, and Chlorine

	Atomic radius (pm)	First Ionization Energy (kJ/mol ⁻¹)
Element 1	170	1086.5
Element 2	180	1011.8
Element 3	175	1251.2
Element 4	152	1313.9

- Which element is number 3? Justify your answer using both properties.
- What is the outermost energy level that has electrons in element 2? How many valence electrons does element 2 have?
- Which element would you expect to have the highest electronegativity? Why?
- How many peaks would the PES for element 4 have and what would the relative heights of those peaks be to each other?



4. The above PES belongs to a neutral chlorine atom.
 - (a) What wavelength of light would be required to eject a 3s electron from chlorine?
 - (b) For the PES of a chloride ion, how would the following variables compare to the peaks on the PES above? Justify your answers.
 - (i) Number of peaks
 - (ii) Height of the peaks
5. The photoelectron spectrum of an element is given below:



- (a) Identify the element this spectra most likely belongs to and write out its full electron configuration.
- (b) Using your knowledge of atomic structure, explain the following:
- (i) The reason for the three discrete areas of ionization energies
 - (ii) The justification for there being a total of five peaks
 - (iii) The relative heights of the peaks when compared to one another

CHAPTER 3 ANSWERS AND EXPLANATIONS

Multiple-Choice

1. **D**

This element has five peaks, meaning a total of five subshells. The final peak, which would be located in the $3p$ subshell, is slightly higher than the $3s$ peak to the left of it. A full $3s$ peak has two electrons, therefore, there must be at least three electrons in the $3p$ subshell. The element that best fits this is phosphorus.

2. **B**

The peaks, in order, represent $1s$, $2s$, $2p$, $3s$, and $3p$.

3. **D**

The less ionization energy that is required to remove an electron, the more kinetic energy that electron will have after ejection.

4. **D**

Valence electrons are those in the outermost energy level. In this case, that is the third level, which has five valence electrons in it (two in $3s$ and three in $3p$).

5. **D**

The ion has three more electrons than the neutral atom, meaning the overall repulsion will be greater. The electrons will “push” one another away more effectively, creating a bigger radius.

6. **B**

In all of these transition metals, the $3d$ subshell is the last one to receive electrons (Aufbau's Principle). A $3d$ subshell has 5 orbitals that can hold two electrons each (Pauli's Exclusion Principle). The electrons will enter one by one into each orbital before any pair up (Hund's Rule), meaning manganese will have all five of its $3d$ electrons unpaired.

7. **D**

Moving across a period, atomic size decreases. Therefore, element Z will be farthest to the right (have the most protons), and thus will have the highest electronegativity value.

8. **D**

Potassium's first valence electron is in the fourth energy level, but both chlorine and argon's first valence electron is in the third energy level.

9. **D**

Potassium has the highest number of protons out of all the options, therefore, it exerts a higher nuclear charge on the electrons, pulling them in closer and creating a smaller atomic radius.

10. **B**

Neutral sodium in its ground state has the electron configuration shown in (C). Sodium forms a bond by giving up its one valence electron ($3s^1$) and becoming a positively charged ion with the same electron configuration as neon, which is (B).

11. **D**

As you move from left to right across the periodic table within a single period (from sodium to chlorine), you add protons to the nuclei, which progressively increases the pull of each nucleus on its electrons. So chlorine will have a larger first ionization energy, greater electronegativity, and a smaller atomic radius.

12. **C**

Electrons that are further away from the nucleus require less ionization energy to remove, and thus will have a greater velocity after being ejected. An electron from $3p$ would be further away from the nucleus than electrons in either $1s$ or $2p$. A $4s$ electron would be faster; however, silicon has no electrons in the $4s$ subshell while in its ground state.

13. **A**

Via $c = \nu\lambda$, you can see that there is an inverse relationship between wavelength and frequency (as one goes up, the other goes down). So, ultraviolet radiation has a higher frequency, and via $E = h\nu$, also has more energy.

14. **B**

The atomic mass is the sum of the neutrons and protons in any atom's nucleus. Since atomic number, which indicates the number of protons, is unique to each element we can subtract this from the weight to find the number of neutrons. The atomic number of B is 5; hence, the ^{11}B has 6 neutrons, only 1 in excess. The atomic number of Cl is 17, so ^{37}Cl has 20 neutrons, 3 in excess. The atomic number of Mg is 12, so ^{24}Mg has the same number of neutrons as protons. The element Ga has an atomic number of 31, meaning there are 39 neutrons in the given nucleus.

15. **D**

Mass spectrometry is used to determine the masses for individual atoms of an element. Through mass spectrometry, it is proven that each element has more than one possible mass.

16. **D**

Nonmetals appear on the right side of the periodic table, and so tend to be smaller than the other elements in their period. For this reason, it is easier for them to attract additional electrons, which means they have higher electronegativity values.

17. **B**

The ionization energy will show a large jump when an electron is removed from a full shell. In this case, the jump occurs between the second and third electrons removed, so the element is stable after two electrons are removed. Magnesium (Mg) is the only element on the list with exactly two valence electrons.

18. **C**

Radiation Energy = Binding Energy + Kinetic Energy

Radiation energy (photon) = 2.5 eV + 2.0 eV

Radiation energy (photon) = 4.5 eV

19. **A**

If the wavelength is decreased, that increases the frequency via $c = \lambda\nu$ (remember, the speed of light is constant). An increased frequency increases the amount of photon energy

via $\Delta E = h\nu$. As the binding energy of the electron would not change, the excess radiation energy would turn into kinetic energy, increasing the velocity of the electron.

20. **D**

The intensity of the light is independent of the amount of energy that the light has. Energy is entirely based on frequency and wavelength, and the brightness of the light would not change the amount of radiation energy. Thus, the amount of kinetic energy of the ejected electrons would not change either.

21. **D**

Nitrogen only has two shells of electrons, while phosphorus has three, making nitrogen smaller and more able to attract additional electrons, meaning a higher electronegativity. Nitrogen and oxygen both have two shells, but oxygen has more protons and an effective nuclear charge of +6 vs. nitrogen's effective nuclear charge of +5. Thus, oxygen has a higher electronegativity.

22. **A**

Remember, transition metals lose their s-electrons first when forming an ion. A manganese atom is initially $[\text{Ar}]4s^23d^5$, but the ion becomes $[\text{Ar}]3d^5$, which has a total of five unpaired electrons (as Hund's Rule states, the electrons will remain unpaired as long as there are empty orbitals for them to enter).

Free-Response

1. (a) Ionization energy is the energy required to remove an electron from an atom. The outermost electron in Ca is at the 4s energy level. The outermost electron in Mg is at the

3s level. The outermost electron in Ca is at a higher energy level and is more shielded from the nucleus, making it easier to remove.

- (b) Calcium has two electrons in its outer shell. The second ionization energy will be larger than the first but still comparable because both electrons are being removed from the same energy level. The third electron is much more difficult to remove because it is being removed from a lower energy level, so it will have a much higher ionization energy than the other two.
- (c) The height of the peaks on a PES represent the relative number of electrons in each subshell. In carbon, all three subshells hold two electrons ($1s^2 2s^2 2p^2$), and thus all peaks are the same height. In oxygen, the $2p$ subshell has four electrons, meaning its peak will be twice as high as the other two.
- (d) The valence electron to be removed from magnesium is located in the completed $3s$ subshell, while the electron to be removed from aluminum is the lone electron in the $3p$ subshell. It is easier to remove the electron from the higher-energy $3p$ subshell than from the lower energy (completed) $3s$ subshell, so the first ionization energy is lower for aluminum.
2. (a) The most common mass of a ClO^- ion is 51 amu. $51 \text{ amu} - 16 \text{ amu} = 35 \text{ amu}$, which must be the mass of the most common isotope of chlorine. As mass number is equal to protons + neutrons, and chlorine has 17 protons (its atomic number), $35 - 17 = 18$ neutrons.
- (b) $51(0.75) + 53(0.25) = 51.5 \text{ amu}$

- (c) No. The only subatomic particles that contribute to the mass of any atom are neutrons and protons. Changing the number of electrons does not change the mass significantly.
- (d) An oxygen atom is smaller than a chlorine atom, and as such is more electronegative. The electrons in the bond are thus more attracted to oxygen than chlorine.
3. (a) Element 3 is chlorine. Chlorine and phosphorus would have the largest atomic radii as they both have three energy levels with electrons present. However, chlorine would be smaller than phosphorus because it has more protons (a higher effective nuclear charge). Additionally, chlorine would have a higher ionization energy than phosphorus due to its smaller size and greater number of protons.
- (b) Element 2 is phosphorus, and therefore the outermost energy level would be $n = 3$. Phosphorus has two electrons in $3s$ and three electrons in $3p$, for a total of five valence electrons.
- (c) Electronegativity increases as atomic radius decreases, so it is expected that element 4 (oxygen) would have the highest electronegativity value. Alternatively, electronegativity increases as an energy level comes close to being full, so it is possible that element 3 (chlorine) may have the highest electronegativity, as it is only one electron away from filling its outermost energy level. (Either answer is acceptable with the proper justification.)
- (d) Element 4 is oxygen, so it would be expected to have three peaks in a PES, one for each subshell. The first two peaks would be the same height because there are two electrons each in the $1s$ and $2s$ subshells. The final peak would be twice the height of the others, as there are four electrons in oxygen's $2p$ subshell.

4. (a) The 3s electron belongs to the peak located at 2.44 MJ/mol. First, you need to calculate the amount of energy needed to remove a single 3s electron (rather than a mole of them):

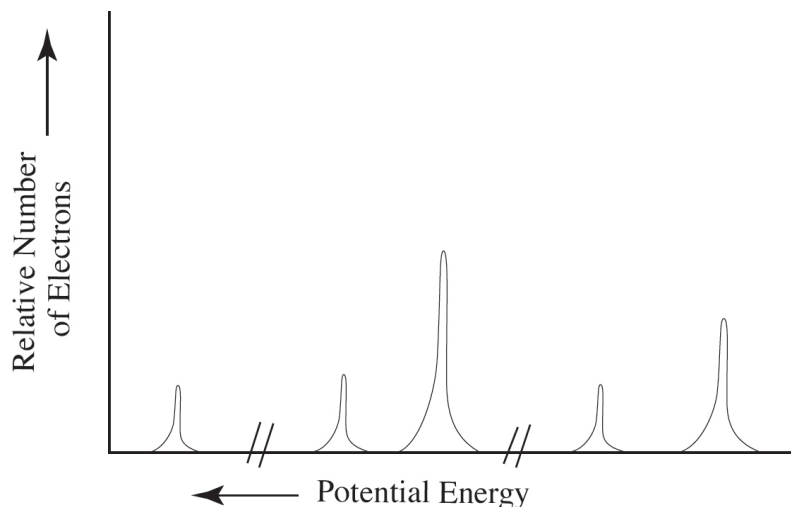
$$\frac{2.44 \text{ MJ}}{1 \text{ mol}} \times \frac{1 \text{ mol}}{6.02 \times 10^{23} \text{ electrons}} = 4.05 \times 10^{-24} \text{ MJ} = 4.05 \times 10^{-18} \text{ J}$$

Then, you can use $E = hc/\lambda$ to calculate the wavelength of light that would have sufficient energy:

$$4.05 \times 10^{-18} \text{ J} = \frac{(6.63 \times 10^{-34} \text{ Js})(3.0 \times 10^8 \text{ m/s})}{\lambda}$$

$$\lambda = 4.9 \times 10^{-8} \text{ m}$$

- (b) (i) The chloride ion would have one more electron, which would enter the 3p energy level. This is represented by the rightmost peak, and so the PES of chloride would have the same number of peaks as that of the chlorine atom.
- (ii) The first four peaks would have the same height, but the last peak would have an additional electron. That would give it six total electrons, making it the same height as the 2p (middle) peak shown in the PES of the neutral chlorine atom.
5. The photoelectron spectrum of an element is given below:



- (a) Identify the element this spectra most likely belongs to and write out its full electron configuration.

This PES belongs to sulfur: $1s^2 2s^2 2p^6 3s^2 3p^4$.

- (b) Using your knowledge of atomic structure, explain the following:

- (i) The reason for the three discrete areas of ionization energies:

Each discrete area of ionization energy represents a different energy level of the electrons. The closer the electrons are to the nucleus, the more ionization energy will be required to remove the electrons. Sulfur has electrons present at three different energy levels, thus, there are three different areas for the peaks.

- (ii) The justification for there being a total of five peaks:

Within each energy level (except for the first), there are subshells which are not the exact same distance from the nucleus. Both energy levels 2 and 3 have *s* and *p* subshells, and while the electrons in those shells will have similar ionization energy values, they will not be

identical. Thus, the five peaks represent $1s$, $2s$, $2p$, $3s$, and $3p$.

- (iii) The relative heights of the peaks when compared to one another:

The heights of the peaks represents the ratio of electrons present in each of them. All three s peaks are exactly one third the height of the $2p$ peak, meaning there are three times more electrons in $2p$ than in any of the s subshells. The $3p$ peak is only twice as high as the s peaks, therefore, there are twice as many electrons in $3p$ than in any of the s subshells.

Chapter 4

Big Idea #2: Bonding and Phases

Chemical and physical properties of materials can be explained by the structure and the arrangement of atoms, ions, or molecules and the forces between them.

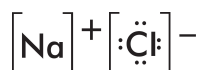
BONDS OVERVIEW

Atoms engage in chemical reactions in order to reach a more stable, lower-energy state. This requires the transfer or sharing of electrons, a process that is called **bonding**. Atoms of elements are usually at their most stable when they have eight electrons in their valence shells. As a result, atoms with too many or too few electrons in their valence shells will find one another and pass the electrons around until all the atoms in the molecule have stable outer shells. Sometimes an atom will give up electrons completely to another atom, forming an ionic bond. Sometimes atoms share electrons, forming covalent bonds.

Ionic Bonds

An ionic solid is held together by the electrostatic attractions between ions that are next to one another in a lattice structure. They often occur between metals and nonmetals. In an ionic bond, electrons are not shared. Instead, the cation gives up an electron (or electrons) to the anion.

The two ions in an ionic bond are held together by electrostatic forces. In the diagram below, a sodium atom has given up its single valence electron to a chlorine atom, which has seven valence electrons and uses the electron to complete its outer shell (with eight). The two atoms are then held together by the positive and negative charges on the ions.



The electrostatic attractions that hold together the ions in the NaCl lattice are very strong, and any substance held together by ionic bonds will usually be a solid at room temperature and have very high melting and boiling points.

Two factors affect the melting points of ionic substances. The primary factor is the charge on the ions. According to Coulomb's Law, a greater charge leads to a greater bond energy (often called lattice energy in ionic bonds), so a compound composed of ions with charges of +2 and -2 (such as MgO) will have a higher melting point than a compound composed of ions with charges of +1 and -1 (such as NaCl). If both compounds are made up of ions with equal charges, then the size of the ions must be considered. Smaller ions will have greater Coulombic attraction (remember, size is inversely proportional to bond energy), so a substance like LiF would have a greater melting point than KBr.

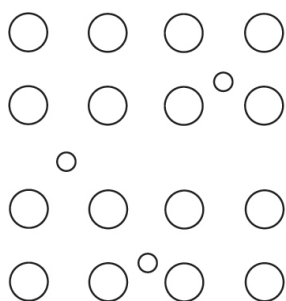
In an ionic solid, each electron is localized around a particular atom, so electrons do not move around the lattice; this makes ionic solids poor conductors of electricity. Ionic liquids, however, do conduct electricity because the ions themselves are free to move about in the liquid phase, although the electrons are still localized around particular atoms. Salts are held together by ionic bonds.

Metallic Bonds

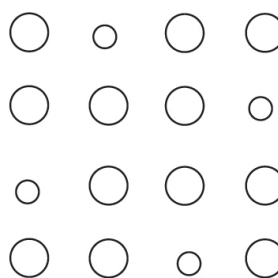
When examining metals, the sea of electrons model can be used. The positively charged core of a metal, consisting of its nucleus and core electrons, is generally stationary, while the valence electrons on each atom do not belong to a specific atom and are very mobile. These mobile electrons explain why metals are such good conductors of electricity. The delocalized structure of a metal also explains why metals are both malleable and ductile, as deforming the metal does not change the environment immediately surrounding the metal cores.

Metals can also bond with each other to form alloys. This typically occurs when two metals are melted into their liquid phases, and are then poured together before cooling and creating the alloy. In an **interstitial alloy**, metal atoms with two vastly different radii combine. Steel is one such example—the much smaller carbon atoms occupy the interstices of the iron atoms. A **substitutional alloy**

forms between atoms of similar radii. Brass is a good example, atoms of zinc are substituted for some copper atoms to create the alloy.



Interstitial
Alloy



Substitutional
Alloy

Covalent Bonds

In a covalent bond, two atoms share electrons. Each atom counts the shared electrons as part of its valence shell. In this way, both atoms achieve complete outer shells.

In the diagram below, two fluorine atoms, each of which has seven valence electrons and needs one electron to complete its valence shell, form a covalent bond. Each atom donates an electron to the bond, which is considered to be part of the valence shell of both atoms.



The first covalent bond formed between two atoms is called a sigma (σ) bond. All single bonds are sigma bonds. If additional bonds between the two atoms are formed, they are called pi (π) bonds. The second bond in a double bond is a pi bond and the second and third bonds in a triple bond are also pi bonds. Double and triple bonds are stronger and shorter than single bonds, but they are not twice or triple the strength.

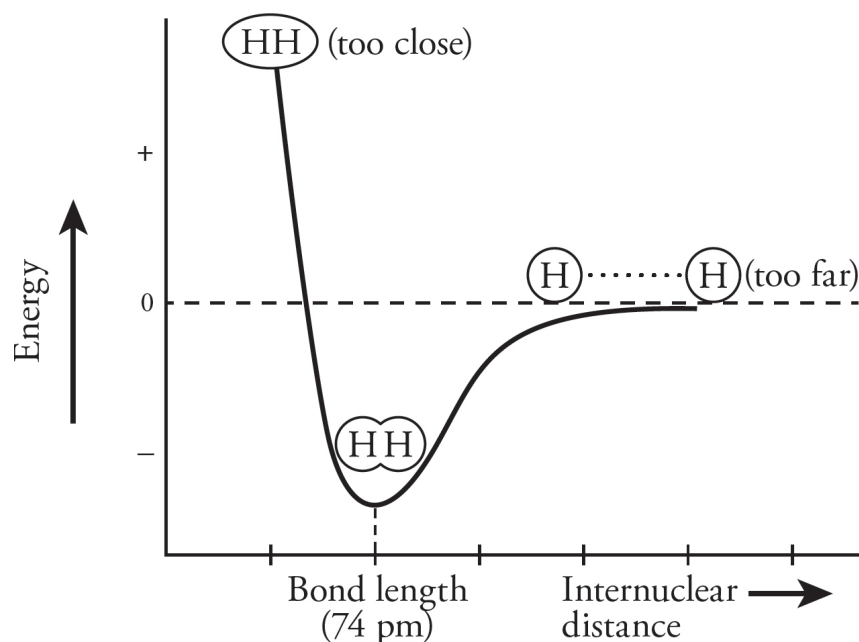
Summary of Multiple Bonds			
Bond type:	Single	Double	Triple
Bond designation:	One sigma (σ)	One sigma (σ) and one pi (π)	One sigma (σ) and two pi (π)
Bond order:	One	Two	Three
Bond length:	Longest	Intermediate	Shortest
Bond energy:	Least	Intermediate	Greatest

Single bonds have one sigma (σ) bond and a bond order of one. The single bond has the longest bond length and the least bond energy.

Internuclear Distance

The length of a covalent bond depends on a balance of two forces. Let's look at a bond between two hydrogen atoms. A bond forms between the hydrogen atoms when the potential energy of that bond is at the minimum possible level. When the hydrogen atoms are too close together, the potential energy value is very high, because the nuclei repel each other. When the atoms are too far apart, the potential energy value is close to zero, because the protons in the nucleus of one hydrogen atom are unable to attract the electrons around the other (and vice-versa).

The minimum potential energy will occur when the repulsive and attractive forces are balanced. A graph of potential energy as a function of the distance between two hydrogen atoms is shown below.



Note that the dashed line marks zero potential energy. However, in this case zero potential energy is not the lowest value. Whenever we are dealing with potential energy, the more negative that value is, the lower the energy is. This is the same concept as looking at the potential energy of electrons at varying distances from the nucleus, as we did on [this page](#).

In this case, the bond between the hydrogen atoms will form at the point that is lowest on the curve. That represents a distance of 74 picometers. The bond length of any covalent bond will always occur at the point of minimum potential energy.

Network (Covalent) Bonds

In a network solid, atoms are held together in a lattice of covalent bonds. You can visualize a network solid as one big molecule. Network solids are very hard and have very high melting and boiling points.

The electrons in a network solid are localized in covalent bonds between particular atoms, so they are not free to move about the lattice. This makes network solids poor conductors of electricity.

The most commonly seen network solids contain either carbon (such as diamond or graphite) and silicon (SiO_2 —quartz). This is because both carbon and silicon have four valence electrons, meaning they are able to form a large number of covalent bonds.

Silicon also serves as a semiconductor when it is doped with other elements. **Doping** is a process in which an impurity is added to an existing lattice. In a normal silicon lattice, each individual silicon atom is bonded to four other silicon atoms. When some silicon atoms are replaced with elements that have only three valence electrons (such as boron or aluminum), the neighboring silicon atoms will lack one bond apiece.

This missing bond (or “hole”) creates a positive charge in the lattice, and the hole attracts other electrons to it, increasing conductivity. Those electrons leave behind holes when they move, creating a chain reaction in which the conductivity of the silicon increases. This type of doping is called **p-doping** for the positively charged holes.

If an element with five valence electrons (such as phosphorus or arsenic) is used to add impurities to a silicon lattice, there is an extra valence electron that is free to move around the lattice, causing an overall negative charge that increases the conductivity of the silicon. This type of doping is called **n-doping** due to the free-moving negatively charged electrons.

LEWIS DOT STRUCTURES



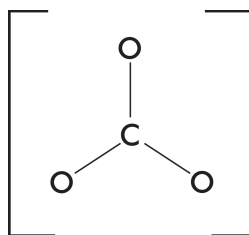
Drawing Lewis Dot Structures

At some point on the test, you'll be asked to draw the Lewis structure for a molecule or polyatomic ion. Here's how to do it.

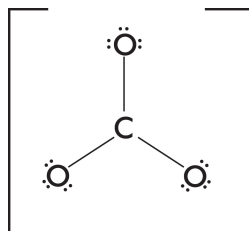
1. Count the valence electrons in the molecule or polyatomic ion; refer to [this page](#) for the periodic table.
2. If a polyatomic ion has a negative charge, add electrons equal to the charge of the total in (1). If a polyatomic ion has a positive charge, subtract electrons equal to the charge of the electrons from the total in (1).
3. Draw the skeletal structure of the molecule and place two electrons (or a single bond) between each pair of bonded atoms. If the molecule contains three or more atoms, the least electronegative atom will usually occupy the central position.
4. Add electrons to the surrounding atoms until each has a complete outer shell.
5. Add the remaining electrons to the central atom.
6. Look at the central atom.
 - (a) If the central atom has fewer than eight electrons, remove an electron pair from an outer atom and add another bond between that outer atom and the central atom. Do this until the central atom has a complete octet.
 - (b) If the central atom has a complete octet, you are finished.
 - (c) If the central atom has more than eight electrons, that's okay too, as long as the total does not exceed twelve.

Let's find the Lewis dot structure for the CO_3^{2-} ion.

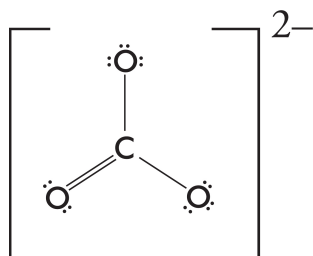
1. Carbon has 4 valence electrons; oxygen has 6.
 $4 + 6 + 6 + 6 = 22$
2. The ion has a charge of -2 , so add 2 electrons.
 $22 + 2 = 24$
3. Carbon is the central atom.



4. Add electrons to the oxygen atoms.

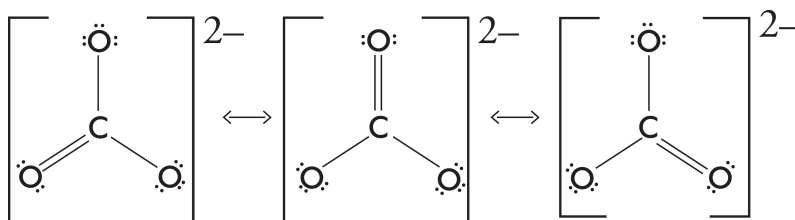


5. We've added all 24 electrons, so there's nothing left to put on the carbon atom.
6. (a) We need to give carbon a complete octet, so we take an electron pair away from one of the oxygens and make a double bond instead. Place a bracket around the model and add a charge of negative two.



Resonance Forms

When we put a double bond into the CO_3^{2-} ion, we place it on any one of the oxygen atoms, as shown below.



The strength and length of all three bonds in the carbonate ion are the same: somewhere between the strength and length of a single bond and a double bond.

To determine the relative length and strength of a bond in a resonance structure, a bond order calculation can be used. A single bond has a bond order of 1, and a double bond has an order of 2. When resonance occurs, pick one of the bonds in the resonance structure and add up the total bond order across the resonance forms, then divide that sum by the number of resonance forms.

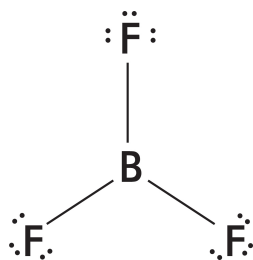
For example, in the carbonate ion above, the top C–O bond would have a bond order of $\frac{1 + 2 + 1}{3}$, or 1.33. Bond order can be used to compare the length and strength of resonance bonds with pure bonds as well as other resonance bonds.

Don't Be Fooled By Appearances

Despite the conventional way the Lewis diagrams have been drawn, remember that the bonds of a compound that displays resonance are all identical. The only reason it looks like a mix of single and double bonds is because any attempt to illustrate a bond somewhere between these two strengths and lengths would be awkward.

Incomplete Octets

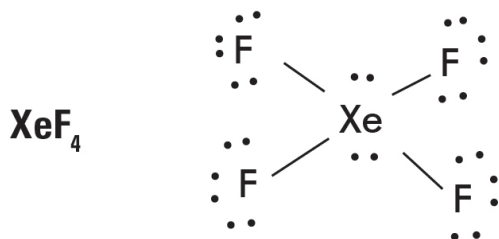
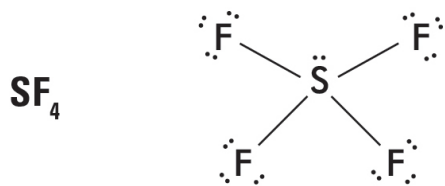
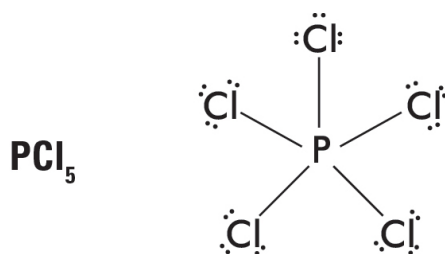
Some atoms are stable with less than eight electrons in their outer shell. Hydrogen only requires two electrons, as does helium (although helium never forms bonds). Boron is considered to be stable with six electrons, as in the BF_3 diagram below. All other atoms involved in covalent bonding require a minimum of eight electrons to be considered stable.



Expanded Octets

In molecules that have *d* subshells available, the central atom can have more than eight valence electrons, but never more than twelve. This means any atom of an element from $n = 3$ or greater can have expanded octets, but NEVER elements in $n = 2$ (C, N, O, etc.). Expanded octets also explains why some noble gases can actually form bonds; the extra electrons go into the empty *d*-orbital.

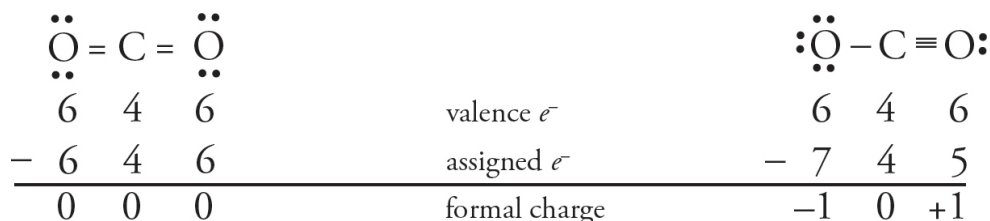
Here are some examples.



Formal Charge

Sometimes, there is more than one valid Lewis structure for a molecule. Take CO₂; it has two valid structures as shown below. To

determine the more likely structure, formal charge is used. To calculate the formal charge on atoms in a molecule, take the number of valence electrons for that atom and subtract the number of assigned electrons in the Lewis structure. When counting assigned electrons, lone pairs count as two and bonds count as one.



The total formal charge for a neutral molecule should be zero, which it is on both diagrams. Additionally, the fewer number of atoms there are with an actual formal charge, the more likely the structure will be —so the left structure is the more likely one for CO_2 . For polyatomic ions, the sum of the formal charges on each atom should equal the overall charge on the ion.

Molecular Geometry

Electrons repel one another, so when atoms come together to form a molecule, the molecule will assume the shape that keeps its different electron pairs as far apart as possible. When we predict the geometries of molecules using this idea, we are using the **valence shell electron-pair repulsion (VSEPR) model**.

In a molecule with more than two atoms, the shape of the molecule is determined by the number of electron pairs on the central atom. The central atom forms **hybrid orbitals**, each of which has a standard shape. Variations on the standard shape occur depending on the number of bonding pairs and lone pairs of electrons on the central atom.

Here are some things you should remember when dealing with the VSEPR model.

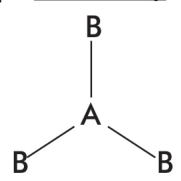
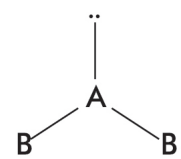
- Double and triple bonds are treated in the same way as single bonds in terms of predicting overall geometry for a molecule; however, multiple bonds have slightly more repulsive strength and will therefore occupy a little more space than single bonds.
- Lone electron pairs have a little more repulsive strength than bonding pairs, so molecules with lone pairs will have slightly reduced bond angles between terminal atoms.

The following pages show the different hybridizations and geometries that you might see on the test.

If the central atom has 2 electron pairs, then it has sp hybridization and its basic shape is **linear**.

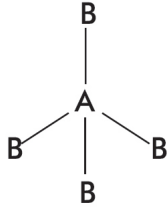
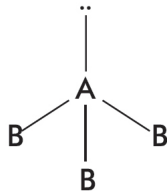
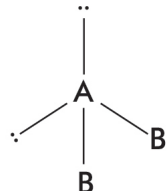
<u>Number of lone pairs</u>	<u>Geometry</u>	<u>Examples</u>
0	B—A—B linear	BeCl ₂ CO ₂

If the central atom has 3 electron pairs, then it has sp^2 hybridization and its basic shape is **trigonal planar**; its bond angles are about 120°.

<u>Number of lone pairs</u>	<u>Geometry</u>	<u>Examples</u>
0	 trigonal planar	BF_3 SO_3 NO_3^- CO_3^{2-}
1	 bent	SO_2

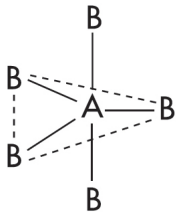
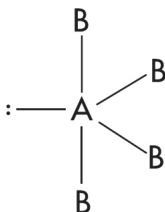
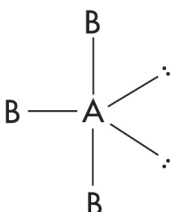
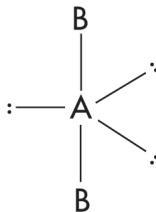
The angle between the terminal atoms in the bent shape is slightly less than 120° because of the extra lone pair repulsion.

If the central atom has 4 electron pairs, then it has sp^3 hybridization and its basic shape is **tetrahedral**; its bond angles are about 109.5° .

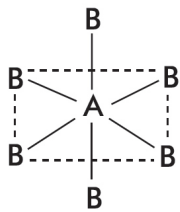
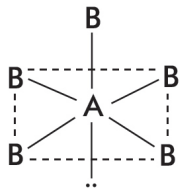
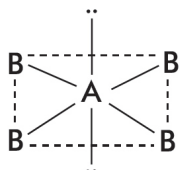
<u>Number of lone pairs</u>	<u>Geometry</u>	<u>Examples</u>
0	 tetrahedral	CH_4 NH_4^+ ClO_4^- SO_4^{2-} PO_4^{3-}
1	 trigonal pyramidal	NH_3 PCl_3 AsH_3 SO_3^{2-}
2	 bent	H_2O OF_2 NH_2^-

The angle between the terminal atoms in the trigonal pyramidal and bent shapes is slightly less than 109.5° because of the extra lone pair repulsion.

If the central atom has 5 electron pairs, its basic shape is **trigonal bipyramidal**.

<u>Number of lone pairs</u>	<u>Geometry</u>	<u>Examples</u>
0	 trigonal bipyramidal	PCl_5 PF_5
1	 folded square, seesaw, distorted tetrahedron	SF_4 IF_4^+
2	 T-shaped	ClF_3 ICl_3
3	 linear	XeF_2 I_3^-

If the central atom has 6 electron pairs, its basic shape is **octahedral**.

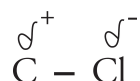
<u>Number of lone pairs</u>	<u>Geometry</u>	<u>Examples</u>
0	 octahedral	SF_6
1	 square pyramidal	BrF_5 IF_5
2	 square planar	XeF_4 ICl_4^-

POLARITY

When a covalent bond forms between two atoms, electrons are shared. However, just because electrons are shared does not mean they are shared equally. For instance, when an atom of fluorine and an atom of carbon bond, the fluorine atom has a stronger pull on the shared electrons due to its higher electronegativity. That means, in the bond between the carbon and the fluorine, the shared electrons (which are constantly moving) will spend more time around the fluorine atom than the carbon one. This type of covalent bond, in which electrons are unequally shared, is called a **polar covalent** bond.

This causes something called dipoles. A **dipole** is an area of either negative or positive charge, and it is based on the likelihood that

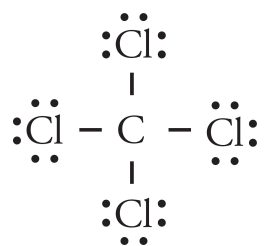
electrons from a bond will be found around an atom. In the diagram below, the carbon has a positive dipole, and the fluorine has a negative dipole. Note that dipoles are indicated by the lowercase version of the Greek letter delta.



Note that if two identical atoms bonds (such as in the Cl_2 molecule), the electrons can be equally shared, creating a **nonpolar covalent** bond in which no dipoles are present.

Molecular Polarity

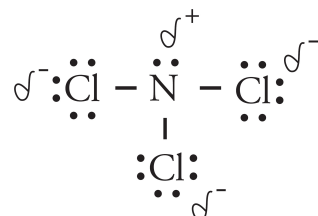
In addition to individual bonds being polar, entire molecules can be polar as well. A molecule's polarity depends on both the polarity of its bonds, but even more importantly, on its overall molecular geometry. Take the carbon tetrachloride molecule, CCl_4 , below:



Even though the C-Cl bond is polar, there are four of them and the molecule is shaped in such a way that the direction of each dipole cancels. Think of it like a giant electron tug-of-war: if each of the four chlorines is pulling on the electrons surrounding carbon with equal strength, all four forces would cancel out directionally, leaving the electrons in the same place. Thus, CCl_4 is a nonpolar molecule.

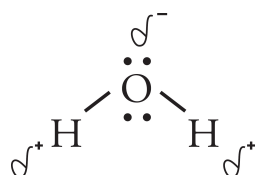
But what if one of those chlorine atoms simply wasn't there? Take NCl_3 , for example. There are still three chlorine atoms pulling on nitrogen's electrons, however, the fourth possible bond location remains unfilled, instead housing a lone pair that belongs entirely to the nitrogen atom. In this case, because chlorine is more

electronegative than nitrogen, each chlorine atom will gain a negative dipole, and the nitrogen atom will gain a positive dipole, as shown below.

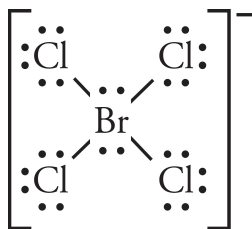


Whether or not a molecule is polar is highly dependent on its molecular geometry. As a general rule of thumb, if the molecule is made up of different atoms and there are any lone pairs on the central atom, the resulting molecule is asymmetrical and will be polar. If there are no lone pairs on the central atom, the molecule is symmetrical and the resulting molecule will be nonpolar.

Also, because Lewis diagrams are specifically drawn with the LEAST electronegative atom in the center (that is, the atom least likely to attract electrons), in polar molecules the central atom generally has a positive dipole, while the terminal atoms have negative dipoles. An exception to this is molecules that have hydrogen as a terminal atom. Hydrogen's electronegativity value is low enough that it will always have a positive dipole in a polar molecule, such as in the water molecules below:



Finally, one last note—one common exception of the “lone pairs on the central atom make things polar” is the square planar shape, such as BrCl_4^- below. The four terminal atoms in a square planar shape are all in the same plane, and as such, square planar molecules are frequently nonpolar.



The term “exception” sure did get used a lot in this section, but when it comes down to it, understanding polarity is about combining the concepts of electronegativity and molecular geometry and figuring out the end result. While the above section gives some good quick-and-dirty tips on figuring out molecular polarity, the best way of learning this concept is to fully understand how those two concepts interact and then applying that knowledge.

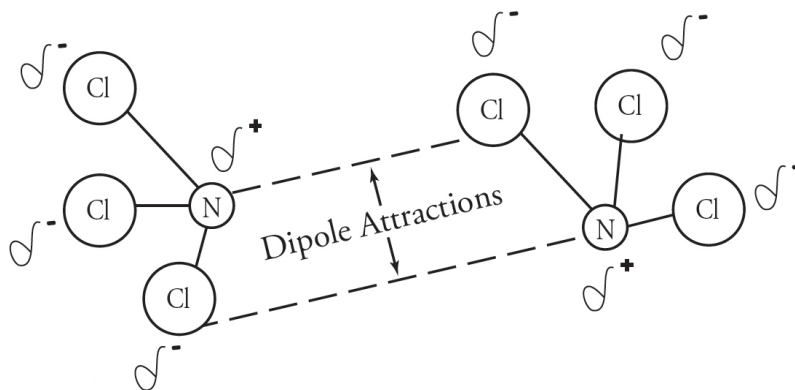
INTERMOLECULAR FORCES

Intermolecular forces (IMFs) are the forces that exist between molecules in a covalently bonded substance. These forces are what need to be broken apart in order for covalent substances to change phases. Note that when ionic substances change phase, bonds between the individual ions are actually broken. When covalent substances change phase, the bonds between the individual atoms remain in place, it is just the forces that hold the molecules to other molecules that break apart.

Dipole–Dipole Forces

Dipole–dipole forces occur between polar molecules: The positive end of one polar molecule is attracted to the negative end of another polar molecule.

Molecules with greater polarity will have greater dipole–dipole attraction, so molecules with larger dipole moments tend to have higher melting and boiling points. Dipole–dipole attractions are relatively weak, however, and these substances melt and boil at very low temperatures. Most substances held together by dipole–dipole attraction are gases or liquids at room temperature.

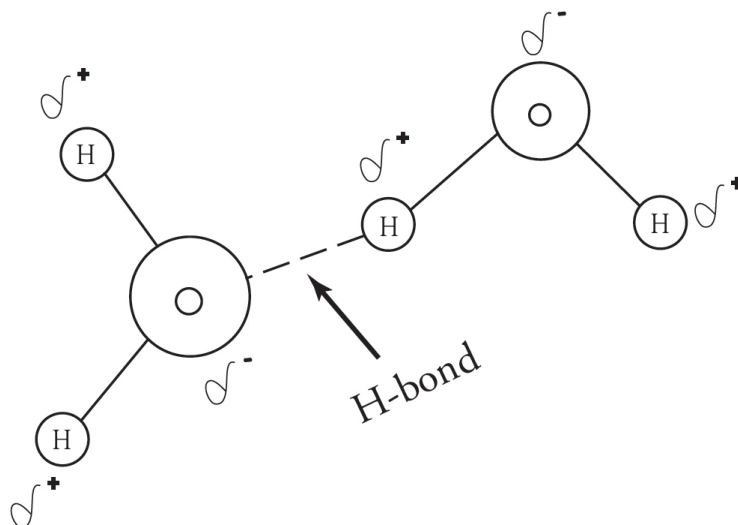


Hydrogen Bonds

Hydrogen bonds are a special type of dipole–dipole attraction. In a hydrogen bond, the positively charged hydrogen end of a molecule is attracted to the negatively charged end of another molecule containing an extremely electronegative element (fluorine, oxygen, or nitrogen—F, O, N).

Hydrogen bonds are much stronger than normal dipole–dipole forces because when a hydrogen atom gives up its lone electron to a bond, its positively charged nucleus is left virtually unshielded. Substances that have hydrogen bonds, such as water and ammonia, have higher melting and boiling points than substances that are held together only by other types of intermolecular forces.

Water is less dense as a solid than as a liquid because its hydrogen bonds force the molecules in ice to form a crystal structure, which keeps them farther apart than they are in the liquid form.



London Dispersion Forces

London dispersion forces occur between all molecules. These very weak attractions occur because of the random motions of electrons on atoms within molecules. At a given moment, a nonpolar molecule might have more electrons on one side than on the other, giving it an instantaneous polarity. For that fleeting instant, the molecule will act as a very weak dipole.

Since London dispersion forces depend on the random motions of electrons, molecules with more electrons will experience greater London dispersion forces. So among substances that experience only London dispersion forces, the one with more electrons will generally have higher melting and boiling points. London dispersion forces are even weaker than dipole–dipole forces, so substances that experience only London dispersion forces melt and boil at extremely low temperatures and tend to be gases at room temperature.

As molecules gain more electrons, the London dispersion forces between them start to become much more significant. Comparing the boiling point of a nonpolar substance with a large number of electrons vs. a polar substance with less electrons is difficult, and there is no simple rule to follow. For instance, water has hydrogen bonds and a boiling point of 100°C . Butane (C_4H_8) and octane (C_8H_{18}) are both completely nonpolar molecules, and while butane's

boiling point is 34°C , octane's is 125°C . Even though octane has no permanent dipoles, it has so many electrons that its London dispersion forces are significant enough that they create greater intermolecular attractions than even the hydrogen bonds in water.

The role of London dispersion forces is often determined by comparing the molar mass of molecules. However, it is not the mass itself which affects the strength of the IMFs. Rather, it is simply that as mass (based on protons and neutrons) increases, so too do the number of electrons, as the molecule must remain electrically neutral.

IMF Questions: Not So Impossible

The AP Exam will most likely focus questions regarding IMFs on molecules with similar masses, as it is much easier to compare the strengths of their IMFs. However, understanding that London dispersion forces can have an impact on IMF-related trends when considering molecules with a large difference in their masses is worth keeping in mind.

Bond Strength

Ionic substances are generally solids at room temperature, and turning them into liquids (melting them), requires the bonds holding the lattice together to be broken. The amount of energy needed for that is based on the Coulombic attraction between the molecules.

Covalent substances, which are liquid at room temperature, will boil when the intermolecular forces between them are broken. For molecules with similar sizes, the following IMF ranking can help you determine the relative strength of the IMFs within the molecules.

- a. Hydrogen bonds
- b. Non-hydrogen bond permanent dipoles
- c. London dispersion forces (temporary dipoles)

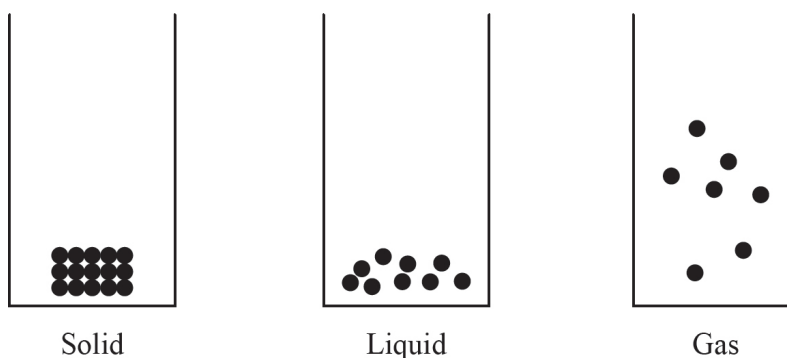
- i. Larger molecules are more polarizable and have stronger London dispersion forces because they have more electrons.

The melting and boiling points of covalent substances are almost always lower than the melting and boiling point of ionic ones.

Metallic bonding, which often only involves one type of atom, tends to be very strong and thus metals (particularly the transition metals) tend to have high melting points. Network covalent bonding is the strongest type of bonding there is, and it is very difficult to cause substances that exhibit network covalent bonding to melt.

Bonding and Phases

The phase of a substance is directly related to the strength of its intermolecular forces. Solids have highly ordered structures where the atoms are packed tightly together, while gases have atoms spread so far apart that most of the volume is free space.



In other words, substances that exhibit weak intermolecular forces (such as London Dispersion forces) tend to be gases at room temperature. Nitrogen (N_2) is an example of this. Substances that exhibit strong intermolecular forces (such as hydrogen bonds) tend to be liquids at room temperature. A good example is water.

Ionic substances do not experience intermolecular forces. Instead, their phase is determined by the ionic bond holding the ions together in the lattice. Because ionic bonds are generally significantly stronger

than intermolecular forces in covalent molecules, ionic substances are usually solid at room temperatures.

VAPOR PRESSURE

Beyond helping to determine the melting point and boiling point of covalent substances, the relative strength of the intermolecular forces in a substance can also predict several other properties of that substance. The most important of these is vapor pressure. Vapor pressure arises from the fact that the molecules inside a liquid are in constant motion. If those molecules hit the surface of the liquid with enough kinetic energy, they can escape the intermolecular forces holding them to the other molecules and transition into the gas phase.

This process is called vaporization. It is not to be confused with a liquid boiling. When a liquid boils, energy (in the form of heat) is added, increasing the kinetic energy of all of the molecules in the liquid until all of the intermolecular forces are broken. For vaporization to occur, no outside energy needs to be added. Note that there is a direct relationship between temperature and vapor pressure. The higher the temperature of a liquid, the faster the molecules are moving and the more likely they are to break free of the other molecules. So, temperature and vapor pressure are directly proportional.

If two liquids are at the same temperature, the vapor pressure is dependent primarily on the strength of the intermolecular forces within that liquid. The stronger those intermolecular forces are, the less likely it is that molecules will be able to escape the liquid, and the lower the vapor pressure for that liquid will be.

SOLUTION SEPARATION

You can use intermolecular forces and the various Coulombic attractions that occur between ions and polar molecules in order to

help separate various substances out from each other. There are several ways to do this.

Solutes and Solvents

There is a basic rule for remembering which solutes will dissolve in which solvents.

Like dissolves like.

That means that polar or ionic solutes (such as salt) will dissolve in polar solvents (such as water). That also means that nonpolar solutes (such as oils) are best dissolved in nonpolar solvents. When an ionic substance dissolves, it breaks up into ions. That's **dissociation**. Free ions in a solution are called electrolytes because they can conduct electricity.

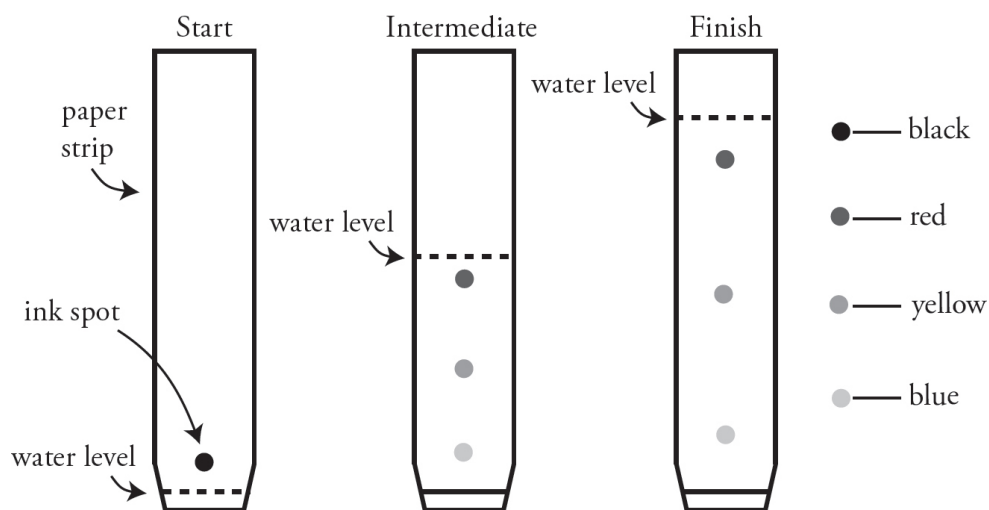
The more ions that are present in an ionic compound, the greater the conductivity of that compound will be when those ions are dissociated. For instance, a solution of magnesium chloride will dissociate into three ions (one Mg^{2+} and two Cl^-). A solution of sodium chloride will dissociate into just two ions (Na^+ and Cl^-). Thus, a solution of magnesium chloride will conduct electricity better than a solution of sodium chloride if both solutions have identical concentrations.

Paper Chromatography

Chromatography is the separation of a mixture by passing it in solution through a medium in which the components of the solution move at different rates. There are several major types of chromatography. The first is paper chromatography, in which paper is the medium through which the solution passes.

Many chemical solutions, such as the ink found in most pens, are a mixture of a number of covalent substances. Each of these substances has its own polarity value, and thus has a different affinity depending on the solvent. One of the most common paper chromatography experiments involves the separation of pigments in black ink. Black ink is usually made up of substances of several different colors, which when combined create black.

In paper chromatography, a piece of filter paper is suspended above a solvent so that the very bottom of the paper is touching the solvent. The ink in question is dotted onto a line at the bottom of the filter paper that starts out just above the solvent level. As the solvent climbs the paper, the various substances inside the ink will be attracted to the polar water molecules. The more polar the substance is, the more it will be attracted to the water molecules, and the further it will travel. You might end up with something that looks like this:



Looking at that strip, you can conclude that the ink was made of three different substances. The one that traveled the farthest with the water (the red pigment) experienced the strongest attractions and was the most polar, whereas the one that didn't travel very far from the original starting line (the blue pigment) was the least polar. Paper chromatography is the most useful with colored substances, which is why ink is used in the above example. If there were

components to the ink that had no visible color, you would not be able to see them on the filter paper, and that is one major limitation of paper chromatography.

The distance the ink travels along the paper is measured via the retention (or retardation) factor, also known as the R_f value. The R_f value is calculated as such:

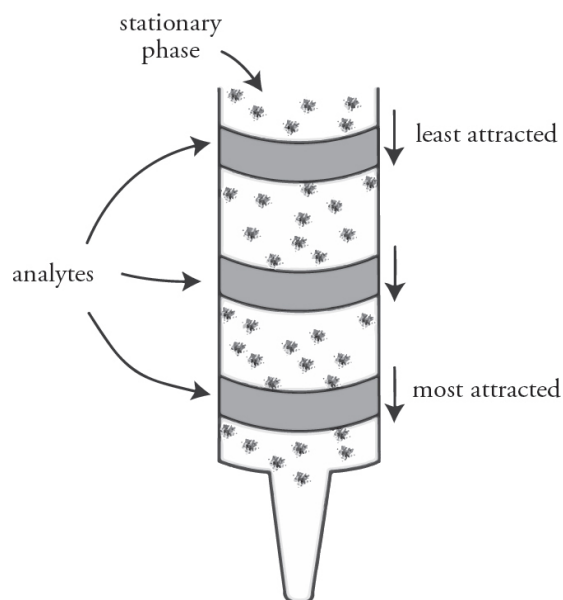
$$R_f = \frac{\text{Distance traveled by solute}}{\text{Distance traveled by solvent front}}$$

The stronger the attraction between the solute and the solvent front is, the larger the R_f value will be. In the diagram above, the red pigment would have the highest R_f value.

Water is not the only solvent that can be used in polar chromatography. There are many nonpolar solvents (such as cyclohexane) that can be used instead. In the case of a nonpolar solvent, the position of the various ink components in the above diagram would have been reversed—the most nonpolar substance would travel the furthest, and the most polar substance would travel the least.

Column Chromatography

Another type of chromatography is column chromatography. In this process, a column is packed with a stationary substance. Then, the solution to be separated (the analyte) is injected into the column, where it adheres to the stationary phase. After that, another solution (called the eluent) is injected into the column. As the eluent passes through the stationary phase, the analyte molecules will be attracted to it with varying degrees of strength depending on their polarity. The more attracted certain analyte molecules are to the eluent, the faster they will travel through (“elute”) and leave the column.



The speed at which the substances move through the column can be monitored, and if there is a sufficient polarity difference between the components, they will leave the column at different times, allowing them to be separated. Generally, after collecting the eluted mixture, it can be analyzed for compositional analysis via a variety of methods.

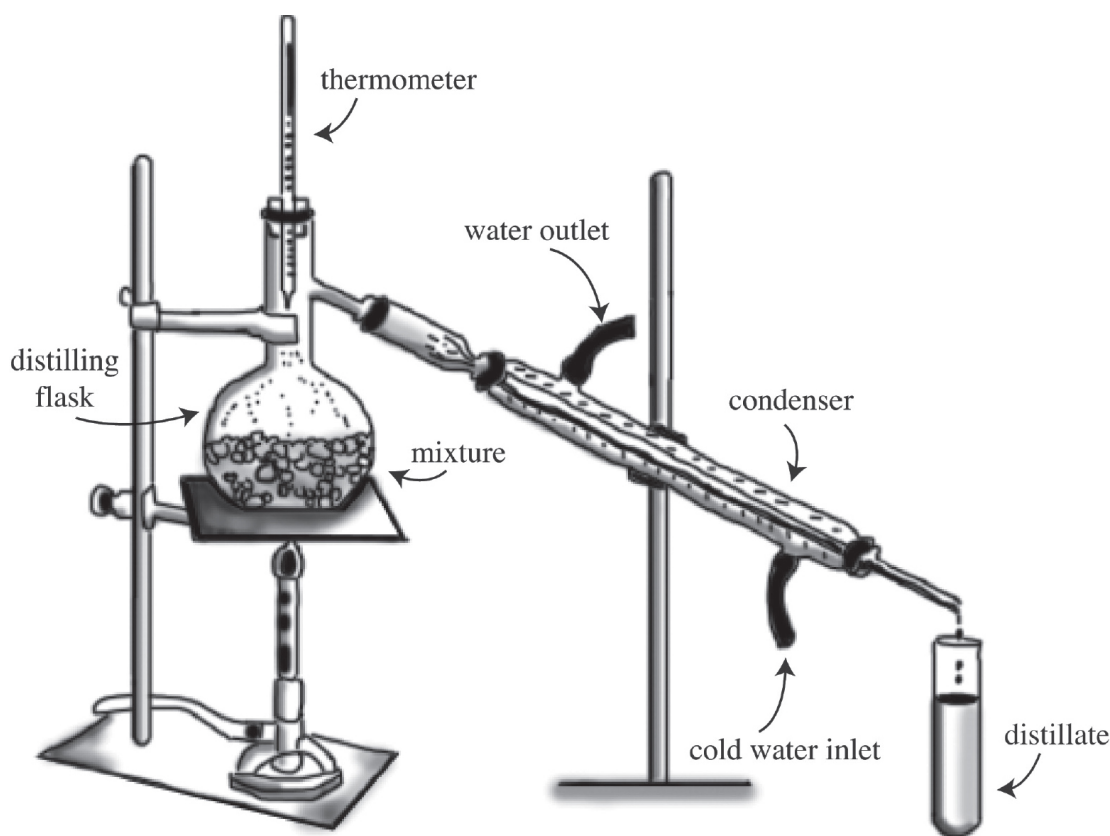
In column chromatography, either liquids or gases can be used as the eluent, depending on the situation.

Distillation

A third method for separating solutions is distillation. Distillation takes advantage of the different boiling points of substances in order to separate them. For instance, if you have a mixture of water (BP: 100°C) and ethanol (BP: 78°C) and then heat that mixture to 85°C , the ethanol will boil but the water will not.

A condenser is a piece of glassware that consists of a smaller tube running through a larger tube. The larger tube has hose connections on it, allowing for water to be run through it. This effectively cools the inner tube and not the outer tube. At that point, the ethanol vapor, when run through the inner tube, will cool and condense back

into a liquid form, which can be collected on the other side of the condenser.



A major advantage to distillation is that the solutions need not be colored at all to separate them. Keeping the flask at a constant temperature can be a challenge, which is why the temperature must be monitored closely to ensure that you are only boiling one component of the mixture at a time. The biggest disadvantage is that it cannot be used to separate a mixture that contains substances with unknown boiling points.

KINETIC MOLECULAR THEORY

For ideal gases, the following assumptions can be made:

- The kinetic energy of an ideal gas is directly proportional to its absolute temperature: The greater the temperature, the greater the average kinetic energy of the gas molecules.

The Average Kinetic Energy of a Single Gas Molecule

$$KE = \frac{1}{2}mv^2$$

m = mass of the molecule (kg)

v = speed of the molecule (meters/sec)

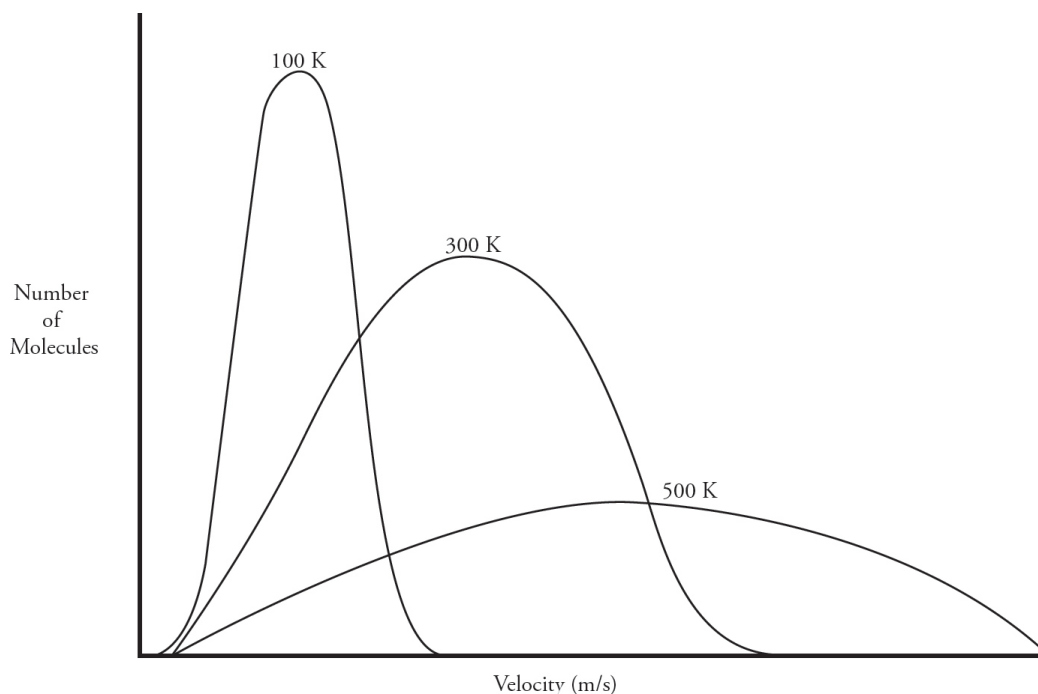
KE is measured in joules

- If several different gases are present in a sample at a given temperature, all the gases will have the same average kinetic energy. That is, the average kinetic energy of a gas depends only on the absolute temperature, not on the identity of the gas.
- The volume of an ideal gas particle is insignificant when compared with the volume in which the gas is contained.
- There are no forces of attraction between the gas molecules in an ideal gas.
- Gas molecules are in constant motion, colliding with one another and with the walls of their container without losing any energy.

MAXWELL-BOLTZMANN DIAGRAMS

A Maxwell-Boltzmann diagram shows the range of velocities for molecules of a gas. Molecules at a given temperature are not all moving at the same velocity. When determining the temperature, we take the average velocity of all the molecules and use that in the relevant equation to calculate temperature. You do not need to know that equation (unless you are taking AP Physics!). All you need to know here is that temperature is directly proportional to kinetic energy.

The first type of Maxwell-Boltzmann diagram involves plotting the velocity distributions for the molecules of one particular gas at multiple temperatures. In the diagram below, there are three curves representing a sample of nitrogen gas at 100 K, 300 K, and 500 K.



As you can see, the higher the temperature of the gas, the larger the range is for the velocities of the individual molecules. Gases at higher temperatures have greater kinetic energy (KE), and as all the molecules in this example have the same mass, the increased KE is due to the increased velocity of the gas molecules.



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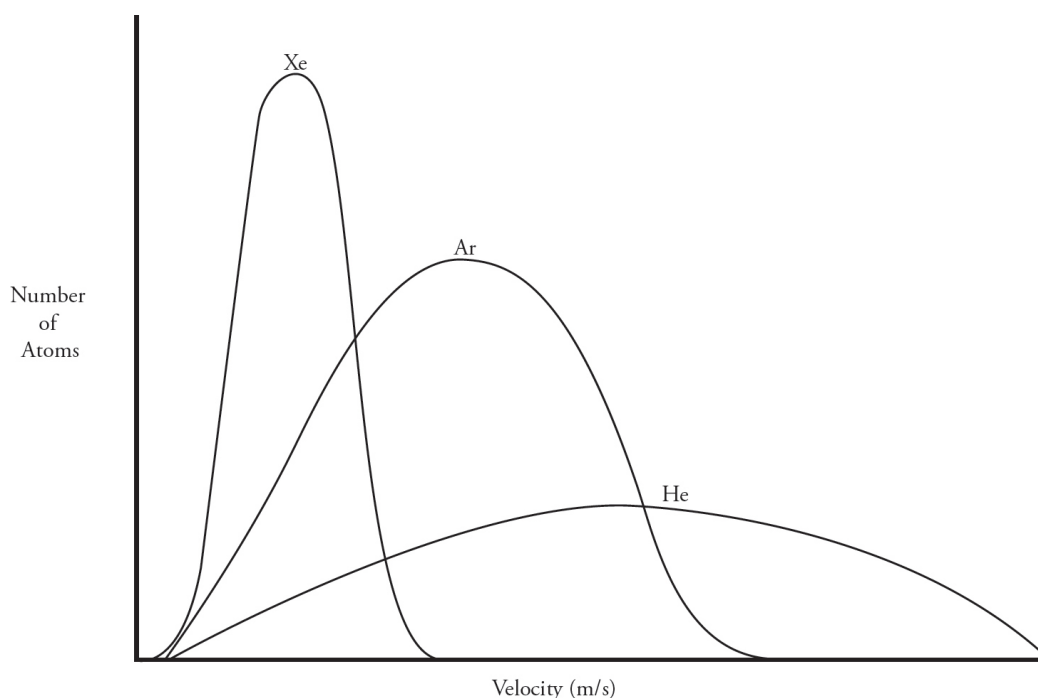
If you wish to dig deeper into many of these topics, check out these other books from The Princeton Review:

Cracking AP Physics 1

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Maxwell-Boltzmann diagrams are also used to show a number of different gases at the same temperature. The diagram on the next page shows helium, argon, and xenon gas, all at 300 K:



In this case, all of the gases have the same amount of total kinetic energy because they have identical temperatures. However, not all of the atoms have the same mass. If the atoms have smaller masses, they must have greater velocities in order to have a kinetic energy

identical to that of atoms with greater mass. Because helium atoms have the least mass, they have the highest average velocity. Xenon atoms, which have a much greater mass, have a correspondingly lower velocity.

EFFUSION

Effusion is the rate at which a gas will escape from a container through microscopic holes in the surface of the container. For instance, even though the rubber or latex that makes up a balloon may seem solid, after the balloon is filled with a gas, it will gradually shrink over time. This is due to the fact that there are tiny holes in the surface of the balloon, through which the even tinier gas molecules can escape.

The rate at which a gas effuses from a container is dependent on the speed of the gas particles. The faster the particles are moving, the more often they hit the sides of the container, and the more likely they are to hit a hole and escape. The rate of effusion thus increases with temperature, but also, if examining gases at the same temperature, the gas with the lower molar mass will effuse first.

It is likely you have experienced this; a balloon filled with helium will deflate more rapidly than one filled with air (which is composed primarily of nitrogen and oxygen) or one filled with carbon dioxide. There is a formula that quantifies the rate at which a gas will effuse, but that is beyond the scope of the exam. As long as you understand the basic principles behind effusion, you should be able to answer any questions on this topic that may come up.

THE IDEAL GAS EQUATION

You can use the ideal gas equation to calculate any of the four variables relating to the gas, provided that you already know the other three.

The Ideal Gas Equation

$$PV = nRT$$

P = the pressure of the gas (atm)

V = the volume of the gas (L)

n = the number of moles of gas

R = the gas constant, 0.0821 L·atm/mol·K

T = the absolute temperature of the gas (K)

You can also manipulate the ideal gas equation to figure out how changes in each of its variables affect the other variables. The following equation, often called the Combined Gas Law, can only be used when the number of moles is held constant.

$$\frac{P_1V_1}{T_1} = \frac{P_2V_2}{T_2}$$

P = the pressure of the gas (atm)

V = the volume of the gas (L)

T = the absolute temperature of the gas (K)

You should be comfortable with the following simple relationships:

- If the volume is constant: As pressure increases, temperature increases; as temperature increases, pressure increases.
- If the temperature is constant: As pressure increases, volume decreases; as volume increases, pressure decreases. That's Boyle's Law.

- If the pressure is constant: As temperature increases, volume increases; as volume increases, temperature increases. That's Charles's Law.

DALTON'S LAW

Dalton's Law states that the total pressure of a mixture of gases is just the sum of all the partial pressures of the individual gases in the mixture.

Dalton's Law

$$P_{\text{total}} = P_a + P_b + P_c + \dots$$

You should also note that the partial pressure of a gas is directly proportional to the number of moles of that gas present in the mixture. So if 25 percent of the gas in a mixture is helium, then the partial pressure due to helium will be 25 percent of the total pressure. This concept, often represented by the variable x , is called the mole fraction.

Partial Pressure

$$P_a = (P_{\text{total}})(X_a)$$

$$X_a = \frac{\text{moles of gas A}}{\text{total moles of gas}}$$

DEVIATIONS FROM IDEAL BEHAVIOR

At low temperature and/or high pressure, gases behave in a less-than-ideal manner. That's because the assumptions made in kinetic

molecular theory become invalid under conditions where gas molecules are packed too tightly together.

Two things happen when gas molecules are packed too tightly.

- *The volume of the gas molecules becomes significant.*
The ideal gas equation does not take the volume of gas molecules into account, so the actual volume of a gas under nonideal conditions will be larger than the volume predicted by the ideal gas equation.
- *Gas molecules attract one another and stick together.*
The ideal gas equation assumes that gas molecules never stick together. When a gas is packed tightly together, intermolecular forces become significant, causing some gas molecules to stick together. When gas molecules stick together, there are fewer particles bouncing around and creating pressure, so the real pressure in a nonideal situation will be smaller than the pressure predicted by the ideal gas equation.

Even under normal conditions, gases will still show some deviation from ideal behavior due to the IMFs between the various gas molecules. These interactions are generally minimal, but when considering the likelihood that a gas will deviate from ideal behavior, stronger IMFs will lead to more deviations. H_2O , with hydrogen bonding, would be significantly more likely to deviate from ideal behavior than CH_4 , which only has LDFs.

If you are looking at gases that have similar IMF types, remember that the more electrons a gas has the more polarizable it is, and is more likely to deviate from ideal behavior. For instance, within the noble gases, which have only LDFs, argon is more likely to deviate from ideal behavior than helium, but less likely to deviate than xenon.

DENSITY

You may be asked about the density of a gas. The density of a gas is measured in the same way as the density of a liquid or solid: in mass per unit of volume.

Density of a Gas

$$D = \frac{m}{V}$$

D = density

m = mass of gas, usually in grams

V = volume occupied by a gas, usually in liters

The density of any gas sample can also be determined by combining the density equation with the Ideal Gas Law.

If $D = \frac{m}{V}$, then $V = \frac{m}{D}$.

Substituting that into the Ideal Gas Law:

$$\frac{Pm}{D} = nRT$$

A little rearrangement yields:

$$D = \frac{Pm}{nRT}$$

The term (m/n) describes mass per mole, which is how molar mass (MM) is measured. Thus:

$$D = \frac{P(\text{MM})}{RT}$$

If you are given the density of a gas and need to find the molar mass, this can also be rewritten as:

$$MM = \frac{DRT}{P}$$

CHAPTER 4 QUESTIONS

[Click here](#) to download and print the PDF version of this exercise.

Multiple-Choice Questions

1. A sealed, rigid container contains three gases: 28.0 g of nitrogen, 40.0 g of argon, and 36.0 g of water vapor. If the total pressure exerted by the gases is 2.0 atm, what is the partial pressure of the nitrogen?

(A) 0.33 atm
(B) 0.40 atm
(C) 0.50 atm
(D) 2.0 atm
2. Which of the following pairs of elements is most likely to create an interstitial alloy?

(A) Titanium and copper
(B) Aluminum and lead
(C) Silver and tin
(D) Magnesium and calcium
3. Why can a molecule with the structure of NBr_5 not exist?

(A) Nitrogen only has two energy levels and is thus unable to expand its octet.
(B) Bromine is much larger than nitrogen and cannot be a terminal atom in this molecule.
(C) It is impossible to complete the octets for all six atoms using only valence electrons.

- (D) Nitrogen does not have a low enough electronegativity to be the central atom of this molecule.

Use the following information to answer questions 4-7.

An evacuated rigid container is filled with exactly 2.00 g of hydrogen and 10.00 g of neon. The temperature of the gases is held at 0°C and the pressure inside the container is a constant 1.0 atm.

4. What is the mole fraction of neon in the container?
- (A) 0.17
(B) 0.33
(C) 0.67
(D) 0.83
5. What is the volume of the container?
- (A) 11.2 L
(B) 22.4 L
(C) 33.5 L
(D) 48.8 L
6. A sample of liquid NH_3 is brought to its boiling point. Which of the following occurs during the boiling process?
- (A) The N–H bonds within the NH_3 molecules break apart.
(B) The overall temperature of the solution rises as the NH_3 molecules speed up.
(C) The amount of energy within the system remains constant.

(D) The hydrogen bonds holding separate NH_3 molecules together break apart.

7. Which gas particles have a higher RMS velocity and why?

(A) Hydrogen, because it has a lower molar mass

(B) Neon, because it has a higher molar mass

(C) Hydrogen, because it has a larger atomic radius

(D) Neon, because it has a smaller atomic radius

8. Which of the following compounds would have the highest lattice energy?

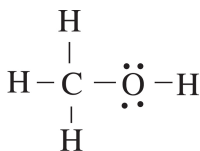
(A) LiF

(B) MgCl_2

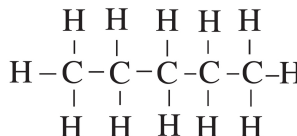
(C) CaBr_2

(D) C_2H_6

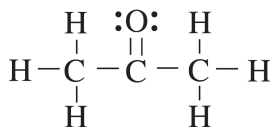
9. The following diagrams show the Lewis structures of four different molecules. Which molecule would travel the farthest in a paper chromatography experiment using a polar solvent?



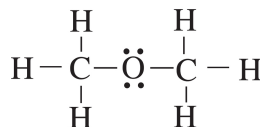
Methanol



Pentane



Acetone



Ether

(A) Methanol

(B) Pentane

(C) Acetone

(D) Ether

10. Which of the following substances has an asymmetrical molecular structure?

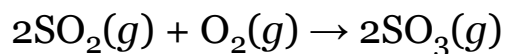
(A) SF_4

(B) PCl_5

(C) BF_3

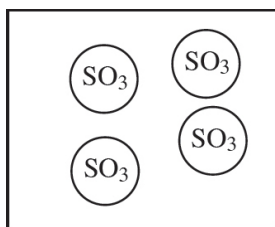
(D) CO_2

Use the following information to answer questions 11-14.

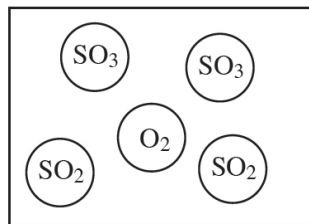


4.0 mol of gaseous SO_2 and 6.0 mol of O_2 gas are allowed to react in a sealed container.

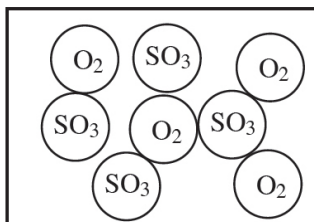
11. Which particulate drawing best represents the contents of the flask after the reaction goes to completion?



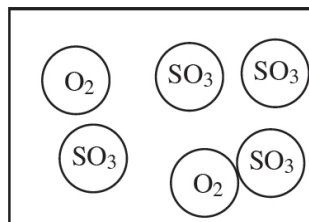
(A)



(B)



(C)



(D)

12. If the temperature remains constant, what percentage of the original pressure will the final pressure in the container be equal to?
- (A) 67%
- (B) 80%
- (C) 100%
- (D) 133%
13. At a given point in the reaction, all three gases are present at the same temperature. Which gas molecules will have the highest velocity and why?
- (A) The O_2 molecules, because they have the least mass
- (B) The O_2 molecules, because they are the smallest
- (C) The SO_3 molecules, because they are products in the reaction
- (D) Molecules of all three gases will have the same speed because they have the same temperature.
14. Under which of the following conditions would the gases in the container most deviate from ideal conditions and why?

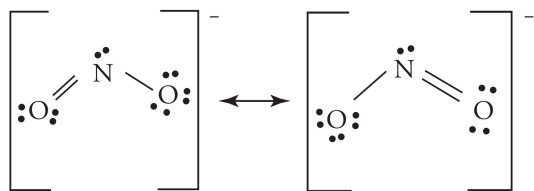
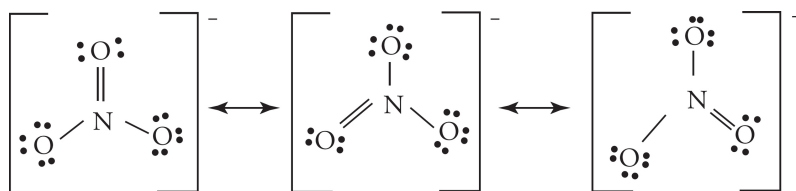
- (A) Low pressures, because the gas molecules would be spread far apart
 - (B) High pressures, because the gas molecules will be colliding frequently
 - (C) Low temperatures, because the intermolecular forces between the gas molecules would increase
 - (D) High temperatures, because the gas molecules are moving too fast to interact with one another
15. A mixture of helium and neon gases has a total pressure of 1.2 atm. If the mixture contains twice as many moles of helium as neon, what is the partial pressure due to neon?
- (A) 0.2 atm
 - (B) 0.3 atm
 - (C) 0.4 atm
 - (D) 0.8 atm
16. Nitrogen gas was collected over water at 25°C. If the vapor pressure of water at 25°C is 23 mmHg, and the total pressure in the container is measured at 781 mmHg, what is the partial pressure of the nitrogen gas?
- (A) 46 mmHg
 - (B) 551 mmHg
 - (C) 735 mmHg
 - (D) 758 mmHg
17. A 22.0 gram sample of an unknown gas occupies 11.2 liters at standard temperature and pressure. Which of the following could be the identity of the gas?
- (A) CO_2

(B) SO_3

(C) O_2

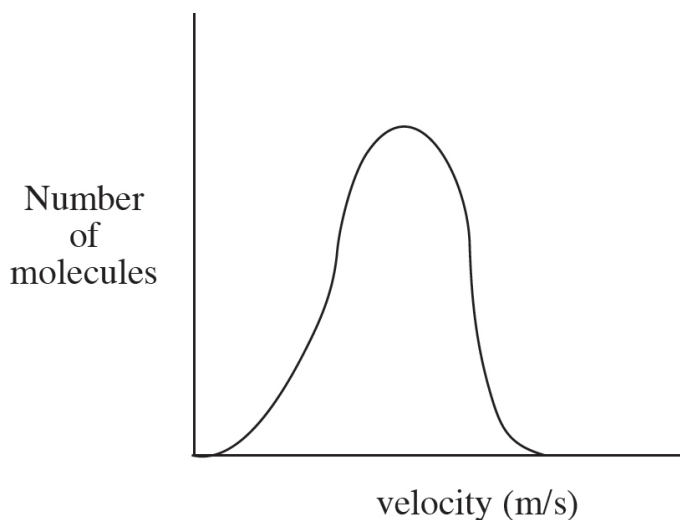
(D) He

18. Lewis diagrams for the nitrate and nitrite ions are shown below. Choose the statement that correctly describes the relationship between the two ions in terms of bond length and bond energy.

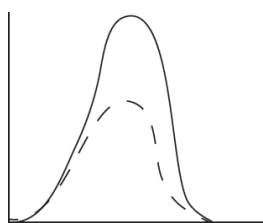


- (A) Nitrite has longer and stronger bonds than nitrate.
(B) Nitrite has longer and weaker bonds than nitrate.
(C) Nitrite has shorter and stronger bonds than nitrate.
(D) Nitrite has shorter and weaker bonds than nitrate.

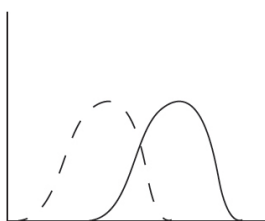
19.



The diagram above shows the speed distribution of molecules in a gas held at 200 K. Which of the following representations would best represent the gas at a higher temperature? (Note: The original line is shown as a dashed line in the answer options.)



(A)



(B)



(C)



(D)

20. A gas sample with a mass of 10 grams occupies 5.0 liters and exerts a pressure of 2.0 atm at a temperature of 26°C. Which of the following expressions is equal to the molecular mass of the gas? The gas constant, R , is 0.08 (L·atm)/(mol·K).

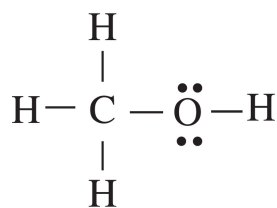
(A) $(0.08)(299) \text{ g/mol}$

(B) $\frac{(299)(0.50)}{(2.0)(0.08)} \text{ g/mol}$

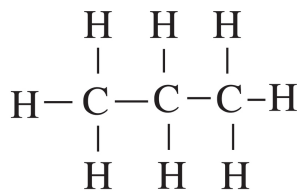
(C) $\frac{299}{0.08} \text{ g/mol}$

(D) $(2.0)(0.08) \text{ g/mol}$

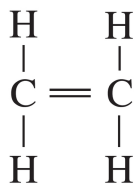
Use the following information to answer questions 21-23.



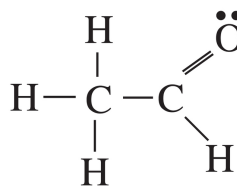
Methanol



Propane



Ethene



Ethanal

21. Based on the strength of the intermolecular forces in each substance, estimate from greatest to smallest the vapor pressures of each substance in liquid state at the same temperature.

- (A) Propane > Ethanal > Ethene > Methanol
(B) Ethene > Propane > Ethanal > Methanol
(C) Ethanal > Methanol > Ethene > Propane
(D) Methanol > Ethanal > Propane > Ethene

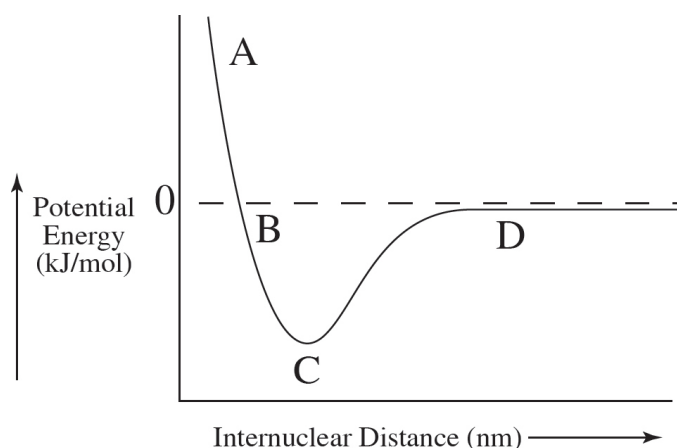
22. When in liquid state, which two substances are most likely to be miscible with water?

- (A) Propane and ethene
- (B) Methanol and propane
- (C) Ethene and ethanal
- (D) Methanol and ethanal

23. Between propane and ethene, which will likely have the higher boiling point and why?

- (A) Propane, because it has a greater molar mass
- (B) Propane, because it has a more polarizable electron cloud
- (C) Ethene, because of the double bond
- (D) Ethene, because it is smaller in size

24. The graph below shows the amount of potential energy between two hydrogen atoms as the distance between them changes. At which point in the graph would a molecule of H_2 be the most stable?



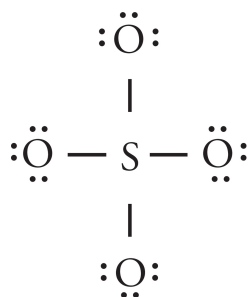
- (A) Point A
- (B) Point B

(C) Point C

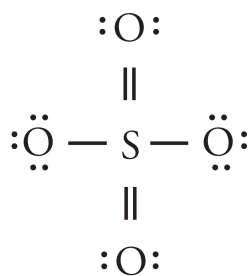
(D) Point D

Use the following information to answer questions 25-28.

There are several different potential different Lewis diagrams for the sulfate ion, two of which are below.



Structure A



Structure B

25. What is the molecular geometry in the structure A?

(A) Tetrahedral

(B) Trigonal Planar

(C) Trigonal Pyramidal

(D) Octahedral

26. What is the S–O bond order in the structure B?

(A) 1.0

(B) 1.33

(C) 1.5

(D) 1.67

27. Which of the following statements regarding the structure B is true?

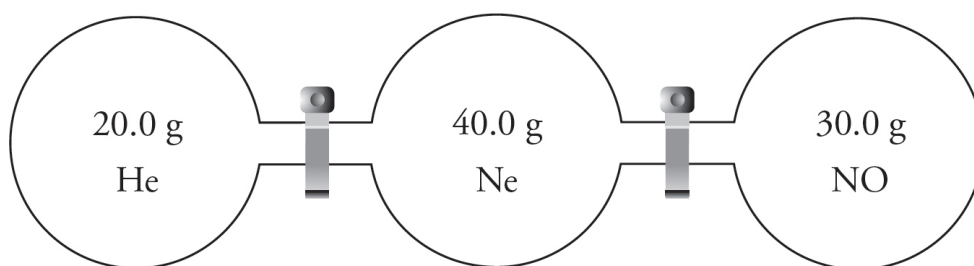
- (A) The double bonds must be located opposite of each other due to additional electron repulsion.
- (B) It is a more polar molecule than the molecule represented by structure A.
- (C) The bonds in the molecule are weaker than those in structure A.
- (D) All bonds in the molecule are identical to each other.

28. Which structure is more likely to correspond with the actual Lewis diagram for the sulfate ion?

- (A) Structure A; single bonds are more stable than double bonds
- (B) Structure A; it has the most unshared pairs of electrons
- (C) Structure B; there are more possible resonance structures
- (D) Structure B; fewer atoms have formal charges

Use the following information to answer questions 29-31.

The diagram below shows three identical 1.0 L containers filled with the indicated amounts of gas. The stopcocks connecting the containers are originally closed and the gases are all at 25°C. Assume ideal behavior.



29. Which gas exerts the greatest pressure?

- (A) He
- (B) Ne

(C) NO

(D) All gases exert the same amount of pressure.

30. Which gas has the strongest IMFs?

(A) He

(B) Ne

(C) NO

(D) All gases have identical IMFs.

31. The stopcocks are opened. If the tubing connecting the containers has negligible volume, by what percentage will the pressure exerted by the neon gas decrease?

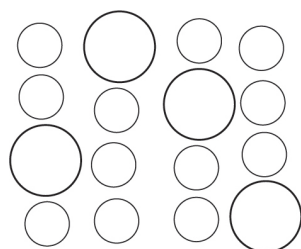
(A) 25%

(B) 33%

(C) 50%

(D) 67%

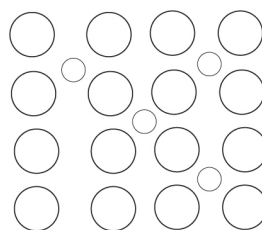
32. Two alloys are shown in the diagrams below—bronze and steel. Which of the following correctly describes the malleability of both alloys compared to their primary metals?



Bronze

○ = Cu

○ = Sn



Steel

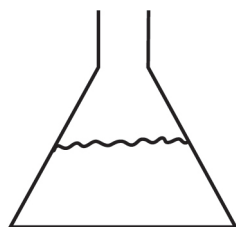
○ = Fe

○ = C

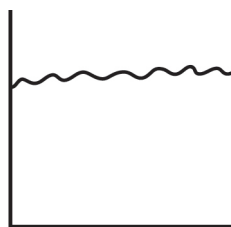
- (A) Bronze's malleability would be comparable to that of copper, but steel's malleability would be significantly lower than that of iron.
- (B) Bronze's malleability would be significantly higher than that of copper, but steel's malleability would be comparable to that of iron.
- (C) Both bronze and steel would have malleability values similar to those of their primary metals.
- (D) Both bronze and steel would have malleability values lower than those of their primary metals.

Free-Response Questions

1. A 250 mL Erlenmeyer flask contains a mixture of two liquids: diethyl ether and ethylamine. The flask is attached to a distillation apparatus and heated until the mixture starts to boil.
 - (a) Why is it important to keep the flask at a constant temperature once the mixture starts to boil?
 - (b)
 - (i) Which liquid is the primary component of the distillate? Justify your answer in terms of IMFs.
 - (ii) The distillate is not a completely pure substance. Why?
 - (c) After the distillate is collected, half of it is transferred into a 250 mL beaker, as shown below. If both containers are left uncovered, which liquid (if either) will evaporate first? Why?



250 ml flask



250 ml beaker

2. The acetyl ion has a formula of $\text{C}_2\text{H}_3\text{O}^-$ and two possible Lewis electron-dot diagram representations:



- (a) Using formal charge, determine which structure is the most likely correct structure.
- (b) For carbon atom “x” in the structure you chose:
- What is the hybridization around the atom?
 - How many sigma and pi bonds has the atom formed?
- (c) A hydrogen ion attaches itself to the acetyl ion, creating $\text{C}_2\text{H}_4\text{O}$. Draw the Lewis diagram of the new molecule.

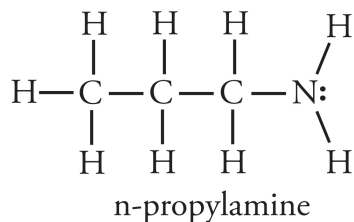
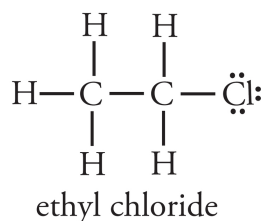
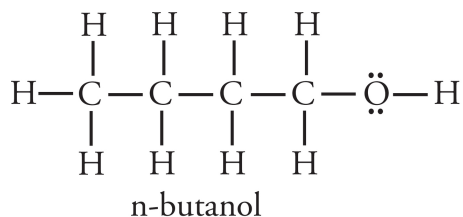
3.

Substance	Boiling Point (C)	Bond Length (Å)	Bond Strength (kcal/mol)
H_2	-253	0.75	104.2
N_2	-196	1.10	226.8
O_2	-182	1.21	118.9
Cl_2	-34	1.99	58.0

- (a) Explain the differences in the properties given in the table above for each of the following pairs.
- The bond strengths of N_2 and O_2
 - The bond lengths of H_2 and Cl_2
 - The boiling points of O_2 and Cl_2

(b) Use the principles of molecular bonding to explain why H_2 and O_2 are gases at room temperature, while H_2O is a liquid at room temperature.

4. A student has a mixture containing three different organic substances. The Lewis diagrams of the substances are below:

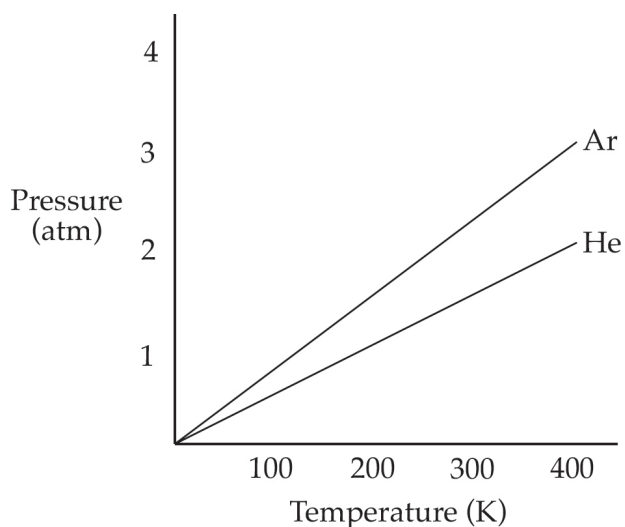


- (a) If the mixture was dabbed onto chromatography paper that was then placed into a nonpolar solvent, rank the R_f values for each component of the mixture from high to low after the solvent has saturated the paper. Justify your answer.
- (b) If the mixture is poured into a chromatography column and then eluted with a very polar substance, which component of the mixture would leave the column first, and why?
- (c) (i) The mixture is heated until it begins to boil. Which substance would be the easiest to separate via distillation, and why?
- (ii) After the substance begins boiling, it continues to be heated at the same rate. Compared to the rate at which it was changing prior to boiling, will the temperature increase faster, slower, or at the same rate? Explain.
- (d) (i) After the components of the mixture have been separated, they are returned to room temperature. Of the

three substances, which would have the highest vapor pressure at room temperature? Justify your answer.

(ii) If the substances were heated (but not boiled), explain what would happen to their vapor pressures.

5.



The graph above shows the changes in pressure with changing temperature of gas samples of helium and argon confined in a closed 2-liter vessel.

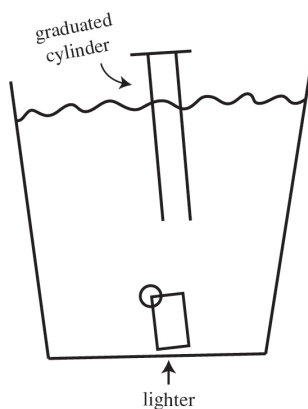
- (a) What is the total pressure of the two gases in the container at a temperature of 200 K?
- (b) How many moles of helium are contained in the vessel?
- (c) How many molecules of helium are contained in the vessel?
- (d) Molecules of which gas will have a greater distribution of velocities at 200 K? Justify your answer.
- (e) If the volume of the container were reduced to 1 liter at a constant temperature of 300 K, what would be the new pressure of the helium gas?

6.



The reaction above took place, and 1.45 liters of oxygen gas were collected over water at a temperature of 29°C and a pressure of 755 millimeters of mercury. The vapor pressure of water at 29°C is 30.0 millimeters of mercury.

- (a) What is the partial pressure of the oxygen gas collected?
 - (b) How many moles of oxygen gas were collected?
 - (c) What would be the dry volume of the oxygen gas at a pressure of 760 millimeters of mercury and a temperature of 273 K?
 - (d) What was the mass of the KClO_3 consumed in the reaction?
7. Equal molar quantities of two gases, O_2 and H_2O , are confined in a closed vessel at constant temperature.
- (a) Which gas, if either, has the greater partial pressure?
 - (b) Which gas, if either, has the greater density?
 - (c) Which gas, if either, has the greater concentration?
 - (d) Which gas, if either, has the greater average kinetic energy?
 - (e) Which gas, if either, will show the greater deviation from ideal behavior?
8. A student performs an experiment in which a butane lighter is held underwater directly beneath a 100-mL graduated cylinder which has been filled with water as shown in the diagram below.



The switch on the lighter is pressed, and butane gas is released into the graduated cylinder. The student's data table for this lab is as follows:

Mass of lighter before gas release	20.432 g
Mass of lighter after gas release	20.296 g
Volume of gas collected	68.40 mL
Water Temperature	19.0°C
Atmospheric Pressure	745 mmHg

- (a) Given that the vapor pressure of water at 19.0°C is 16.5 mmHg, determine the partial pressure of the butane gas collected in atmospheres.
- (b) Calculate the molar mass of butane gas from the experimental data given.
- (c) If the formula of butane is C_4H_{10} , determine the percent error for the student's results.
- (d) The following are common potential error sources that occur during this lab. Explain whether or not each error could have been responsible for the error in the student's results.
 - (i) The lighter was not sufficiently dried before massing it after the gas was released.
 - (ii) The gas in the lighter was not held underwater long enough to sufficiently cool it to the same temperature of the water and was actually at a higher temperature than the water.
 - (iii) Not all of the butane gas released was collected in the graduated cylinder.

CHAPTER 4 ANSWERS AND EXPLANATIONS

Multiple-Choice

1. **C**

There are 1 mole of N_2 , 1 mole of Ar, and 2 moles of water in the container. The mole fraction of nitrogen is: $1/4 = 0.25$.

$$P_{\text{N}_2} = (X_{\text{N}_2})(P_{\text{total}})$$

$$P_{\text{N}_2} = (0.25)(2.0) = 0.50 \text{ atm}$$

2. **B**

Interstitial alloys form when atoms of greatly different sizes combine. The aluminum atoms would have a chance to fit between the comparatively larger lead atoms.

3. **A**

Only atoms with at least three energy levels ($n = 3$ and above) have empty d -orbitals that additional electrons can fit into, thus expanding their octet.

4. **B**

Moles of $\text{H}_2 = \frac{2.00 \text{ g}}{2.00 \text{ g/mol}} = 1.00$ (remember, hydrogen is a diatomic)

$$\text{Moles of Ne} = \frac{10.00 \text{ g}}{20.0 \text{ g/mol}} = 0.500 \text{ mol}$$

Total moles = 1.50 moles.

$$X_{\text{Ne}} = \frac{\text{moles Ne}}{\text{total moles}} = \frac{0.500}{1.500} = 0.33$$

5. **C**

At STP, 1 mole of gas takes up 22.4 L of space: $(1.5)(22.4) = 33.5\text{L}$.

6. **D**

When a covalent substance undergoes a phase change, the bonds between the various molecules inside the substance break apart.

7. **A**

The gas molecules have the same amount of kinetic energy due to their temperature being the same. Via $KE = \frac{1}{2}mv^2$, if KE is the same, then the molecule with less mass must correspondingly have a higher velocity.

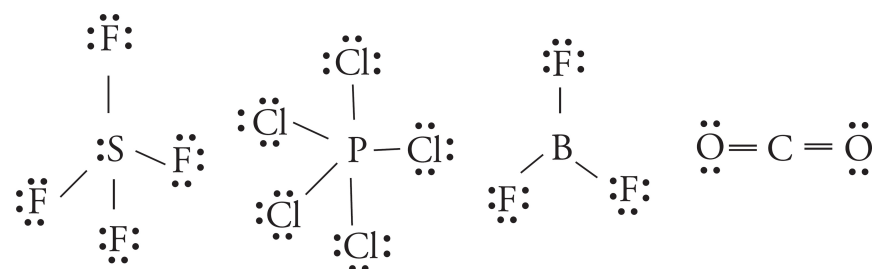
8. **B**

First, C_2H_6 is not an ionic substance and thus has no lattice energy. Next, LiF is composed of ions with charges +1 and -1, and will not be as strong as the two compounds that have ions with charges of +2 and -1. Finally, MgCl_2 is smaller than CaBr_2 , meaning it will have a higher lattice energy, as (according to Coulomb's Law) atomic radius is inversely proportional with bond energy.

9. **A**

In a polar solvent, polar molecules will be the most soluble (like dissolves like). Of the four options, methanol and acetone would both have dipoles, but those of methanol would be significantly stronger due to the H-bonding.

10. **A**



The only tricky bit here is to remember that boron is considered stable with only six electrons in its valence shell.

11. **C**

To determine the number of moles of SO₃ created, stoichiometry must be used.

$$\text{SO}_2: 4 \text{ mol SO}_2 \times \frac{2 \text{ mol SO}_3}{2 \text{ mol SO}_2} = 4 \text{ mol SO}_3$$

$$\text{O}_2: 6 \text{ mol O}_2 \times \frac{2 \text{ mol SO}_3}{1 \text{ mol O}_2} = 12 \text{ mol SO}_3$$

The oxygen is in excess, and only 2.0 mol of it will react. (As every 2.0 mol of SO₂ react with 1 mol of O₂.) Thus, 4.0 mol of SO₃ are created and 4.0 mol of O₂ remain.

12. **B**

If 4.0 mol of SO_3 are created and 4.0 mol of O_2 remain, there are 8.0 mol of gas present after the reaction.

Prior to the reaction there were 10.0 mol of gas. If there are $8/10 = 80\%$ as many moles after the reaction, there is also 80% as much pressure.

13. **A**

If all gases are at the same temperature, they have the same amount of kinetic energy. Given that $KE = \frac{1}{2}mv^2$, if all three gases have the same KE , the gas with the least mass must have the highest velocity in order to compensate.

14. **C**

One of the assumptions of kinetic molecular theory is that the amount of intermolecular forces between the gas molecules is negligible. If the molecules are moving very slowly, as happens when the temperature is lowered, the IMFs between them are more likely to cause deviations from ideal behavior.

15. **C**

From Dalton's Law, the partial pressure of a gas depends on the number of moles of the gas that are present. If the mixture has twice as many moles of helium as neon, then the mixture must be $\frac{1}{3}$ neon. So $\frac{1}{3}$ of the pressure must be due to neon.

$$\left(\frac{1}{3}\right)(1.2 \text{ atm}) = 0.4 \text{ atm.}$$

16. **D**

From Dalton's Law, the partial pressures of nitrogen and water vapor must add up to the total pressure in the container. The partial pressure of water vapor in a closed container will be equal to the vapor pressure of water, so the partial pressure of nitrogen is:

$$781 \text{ mmHg} - 23 \text{ mmHg} = 758 \text{ mmHg}$$

17. **A**

Use the following relationship:

$$\text{Moles} = \frac{\text{liters}}{22.4 \text{ L/mol}}$$

$$\text{Moles of unknown gas} = \frac{11.2 \text{ L}}{22.4 \text{ L/mol}} = 0.500 \text{ moles}$$

$$\text{MW} = \frac{\text{grams}}{\text{mole}}$$

$$\text{MW of unknown gas} = \frac{22.0 \text{ g}}{0.500 \text{ mole}} = 44.0 \text{ grams/mole}$$

That's the molecular weight of CO_2 .

18. **C**

Nitrate has a bond order of $\frac{(1 + 3)}{3} = 1.33$. Nitrite has a bond order of $\frac{(1 + 2)}{2} = 1.5$. A higher bond order means shorter

and stronger bonds.

19. **D**

At a higher temperature, the average velocity of the gas molecules would be greater. Additionally, they would have a greater spread of potential velocities, which would lead to a wider curve.

20. **A**

$MM = \frac{DRT}{P}$. The density of this gas is $\frac{10.0 \text{ g}}{5.0 \text{ L}}$, or $\frac{2.0 \text{ g}}{\text{L}}$.

Plugging that in yields $MM = \frac{(2.0)(0.08)(299)}{(2.0)}$. The value of 2.0 cancels out, leaving the answer (0.08)(299).

21. **B**

Vapor pressure is dependent on intermolecular forces. The weaker the IMFs are, the easier it is for molecules to escape from the surface of the liquid. To begin, polar molecules have stronger IMFs than nonpolar molecules. Methanol and ethanal are both polar, but methanol has hydrogen bonding, meaning it has stronger IMFs (and thus a lower vapor pressure) than ethanal. Ethene and propane are both nonpolar, but propane is larger, meaning it is more polarizable than ethene and thus has stronger IMFs and lower vapor pressure.

22. **D**

Water is polar, and using “like dissolves like,” only polar solvents will be able to fully mix with it to create a homogenous solution.

23. **B**

Both are nonpolar, but propane has a lot more electrons and thus is more polarizable than ethene.

24. **C**

The molecule would be the most stable when it has the largest attractive potential energy, which is represented by a negative sign. While the magnitude of the potential energy may be larger at (A), it is repulsive at that point because the nuclei are too close together.

25. **A**

Four charge clouds and no lone pairs means tetrahedral geometry.

26. **C**

Six total bonds divided by four locations gives a bond order of 1.5.

27. **D**

In any molecule displaying resonance, all bonds are identical.

28. **D**

The formal charge tables for each diagram are below (note: for structure B, the double-bonded oxygens are the first two, and the single bonded are the last two)

Structure A						Structure B					
S	O	O	O	O		S	O	O	O	O	
6	6	6	6	6	Valence e^-	6	6	6	6	6	
4	7	7	7	7	Assigned e^-	6	6	6	7	7	
+2	-1	-1	-1	-1	Formal charge	0	0	0	-1	-1	

The total formal charge on each potential structure is -2 , which is correct, as that is the charge on a sulfate ion. However, the right-hand structure has fewer atoms with formal charges, making it the more likely structure.

29. **A**

Pressure is directly dependent on the number of moles. In their respective containers, there are 5 moles of He, 2 moles of Ne, and 1 mole of NO. As there are the most moles of He, the He must exert the greatest pressure.

30. **D**

One of the precepts of kinetic molecular theory is that gas molecules exert no forces on each other, thus, in all containers there are no IMFs present.

31. **D**

$$P_1 V_1 = P_2 V_2$$

$$P_1(1.0 \text{ L}) = P_2(3.0 \text{ L})$$

$$P_2/P_1 = 1.0/3.0$$

Thus, the pressure of the neon gas is 33% of what it was originally, meaning a decrease of 67%. Note that the same calculation could be used for any of the gases; each gas is expanding to take up three times as much space as it has originally, and thus exerts one-third as much pressure.

32. D

The goal of creating an alloy is often to make the base metal stronger. This means that alloys of any type have less malleability than the metals from which they are created.

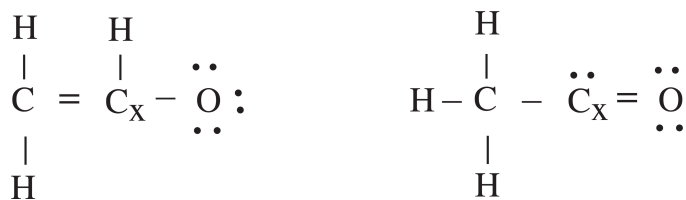
Free-Response

1. (a) If the temperature were to be increased, it may go higher than the boiling point of both substances, not just the one with the lower boiling point. This would create a very impure distillate.

(b) (i) The diethyl ether would be in the distillate. Ethylamine contains hydrogen bonding, which is the strongest type of IMF, while diethyl ether does not.

(ii) Even below the boiling point of ethylamine, some of the ethylamine will spontaneously convert into a gas. The molecules within both liquids are constantly moving, and even if the ethylamine does not boil, some of its molecules will evaporate, convert into a gas, and then be condensed, becoming part of the distillate.

(c) The liquid in the beaker will evaporate first. Evaporation occurs when molecules in the liquid phase have sufficient kinetic energy to break free of the IMFs within the liquid. In order to break free, the molecules must first hit the surface of the liquid. The surface area of the liquid in the beaker is greater than the surface area of the liquid in the test tube, thus, the molecules in the beaker are more likely to hit the surface and evaporate.
2. The acetyl ion has a formula of $\text{C}_2\text{H}_3\text{O}^-$ and two possible Lewis electron-dot diagram representations:



- (a) Using formal charge, determine which structure is the most likely correct and circle it.

For this formal charge calculation, the H atoms are left out as they are identically bonded/drawn in both structures

	C	C _x	O	C	C _x	O		
Valence			4	4	6	4	4	6
Assigned	<u>-4</u>	<u>4</u>	<u>7</u>	<u>-4</u>	<u>5</u>	<u>6</u>		
Formal Charge			0	0	-1	0	-1	0

As oxygen is more electronegative than carbon, an oxygen atom is more likely to have the negative formal charge than a carbon atom. The left-hand structure is most likely correct.

- (b) For carbon atom “x” in the structure you chose:

- (i) What is the hybridization around the atom?

There are three charge groups around the carbon atom, so the hybridization is sp^2 .

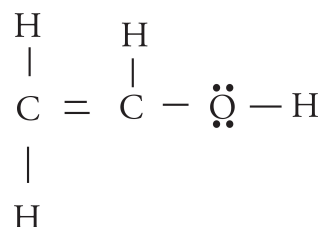
- (ii) How many sigma and pi bonds has the atom formed?

Single bonds consist of sigma bonds, and double bonds consist of one sigma and one pi bond.

There are a total of three sigma bonds and one pi bond around the carbon atom.

- (c) A hydrogen ion attaches itself to the acetyl ion, creating $\text{C}_2\text{H}_4\text{O}$. Draw the Lewis diagram of the new molecule.

The hydrogen ion will attach to the negatively charged oxygen.



3. (a) (i) The bond strength of N_2 is larger than the bond strength of O_2 because N_2 molecules have triple bonds and O_2 molecules have double bonds. Triple bonds are stronger and shorter than double bonds.

(ii) The bond length of H_2 is smaller than the bond length of Cl_2 because hydrogen is a smaller atom than chlorine, allowing the hydrogen nuclei to be closer together.

(iii) Liquid oxygen and liquid chlorine are both nonpolar substances that experience only London dispersion forces of attraction. These forces are greater for Cl_2 because it has more electrons (which makes it more polarizable), so Cl_2 has a higher boiling point than O_2 .

- (b) H_2 and O_2 are both nonpolar molecules that experience only London dispersion forces, which are too weak to form the bonds required for a substance to be liquid at room temperature.

H_2O is a polar substance whose molecules form hydrogen bonds with each other. Hydrogen bonds are strong enough to form the bonds required in a liquid at room temperature.

4. (a) All three components are polar, but ethyl chloride has no hydrogen bonding and thus has the weakest dipoles, meaning it would travel the furthest and have the highest R_f value.

Between n-butanol and n-propylamine, both have H-bonds but the butanol has more electrons, meaning its London dispersion forces are stronger. It thus has the highest polarity and would have the smallest R_f value. So: ethyl chloride > n-propylamine > n-butanol.

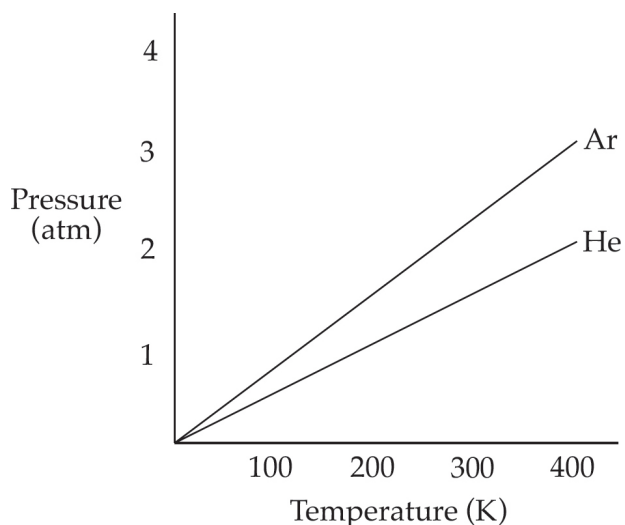
- (b) n-butanol is the most polar and would be most attracted to a polar eluent, and thus would leave the column first.

- (c) (i) Ethyl chloride has the weakest overall IMFs, and thus would have the lowest boiling point and be the easiest to separate out.

- (ii) Prior to boiling, molecules in all three substances were speeding up as heat was added. The increased velocity caused the temperature increase. Once the ethyl chloride starts boiling, though, the heat that would ordinarily be causing the molecules to speed up is instead breaking the IMFs. While the molecules of both the n-butanol and n-propylamine would still be experiencing a velocity increase, those of the ethyl chloride would not. Thus, the overall rate of the temperature change would be less than it was prior to the ethyl chloride starting to boil.

- (d) (i) The substance with the weakest IMFs would allow the largest number of molecules to escape the liquid phase and turn into a gas, which is the cause of vapor pressure. As such, the ethyl chloride should have the highest vapor pressure of the three substances.

- (ii) As temperature increases, so does the molecular velocity. The faster the molecules are going, the more energy they have, and the more likely they are to be able to overcome the IMFs and escape the liquid phase. For all substances, as temperature increases, so does vapor pressure.



5. (a) Read the graph, and add the two pressures.

$$P_{\text{Total}} = P_{\text{He}} + P_{\text{Ar}}$$

$$P_{\text{Total}} = (1 \text{ atm}) + (1.5 \text{ atm}) = 2.5 \text{ atm}$$

- (b) Read the pressure (1 atm) at 200 K, and use the ideal gas equation.

$$n = \frac{PV}{RT} = \frac{(1.0 \text{ atm})(2.0 \text{ L})}{(0.082 \text{ L} \cdot \text{atm}/\text{mol} \cdot \text{K})(200 \text{ K})} = 0.12 \text{ moles}$$

- (c) Use the definition of a mole.

$$\text{Molecules} = (\text{moles})(6.02 \times 10^{23})$$

$$\text{Molecules (atoms) of helium} = (0.12)(6.02 \times 10^{23}) = 7.2 \times 10^{22}$$

(d) As both gases are at the same temperature and thus have the same kinetic energy, the molecules of helium will have a higher average velocity because they have less mass. The higher the average velocity of a gas is, the wider the distribution curve is for the velocities of the individual gas molecules.

(e) Use the following relationship:

$$\frac{P_1 V_1}{T_1} = \frac{P_2 V_2}{T_2}$$

Since T is a constant, the equation becomes:

$$P_1 V_1 = P_2 V_2$$

$$(1.5 \text{ atm})(2.0 \text{ L}) = P_2(1.0 \text{ L})$$

$$P_2 = 3.0 \text{ atm}$$

6. (a) Use Dalton's Law.

$$P_{\text{Total}} = P_{\text{Oxygen}} + P_{\text{Water}}$$

$$(755 \text{ mmHg}) = (P_{\text{Oxygen}}) + (30.0 \text{ mmHg})$$

$$P_{\text{Oxygen}} = 725 \text{ mmHg}$$

(b) Use the Ideal Gas Law. Don't forget to convert to the proper units.

$$n = \frac{PV}{RT} = \frac{\left(\frac{725}{760} \text{ atm}\right)(1.45 \text{ L})}{(0.082 \text{ L} \cdot \text{atm/mol} \cdot \text{K})(302 \text{ K})} = 0.056 \text{ moles}$$

(c) At STP, moles of gas and volume are directly related.

$$\text{Volume} = (\text{moles})(22.4 \text{ L/mol})$$

$$\text{Volume of O}_2 = (0.056 \text{ mol})(22.4 \text{ L/mol}) = 1.25 \text{ L}$$

(d) We know that 0.056 moles of O₂ were produced in the reaction.

From the balanced equation, you know that for every 3 moles of O₂ produced, 2 moles of KClO₃ are consumed. So there are $\frac{2}{3}$ as many moles of KClO₃ as O₂.

$$\text{Moles of KClO}_3 = \left(\frac{2}{3}\right)(\text{moles of O}_2)$$

$$\text{Moles of KClO}_3 = \left(\frac{2}{3}\right)(0.056 \text{ mol}) = 0.037 \text{ moles}$$

$$\text{Grams} = (\text{moles})(\text{MW})$$

$$\text{Grams of KClO}_3 = (0.037 \text{ mol})(122 \text{ g/mol}) = 4.51 \text{ g}$$

7. (a) The partial pressures depend on the number of moles of gas present. Because the number of moles of the two gases are the same, the partial pressures are the same.
- (b) O₂ has the greater density. Density is mass per unit volume. Both gases have the same number of moles in the same volume, but oxygen has heavier molecules, so it has greater density.
- (c) Concentration is moles per volume. Both gases have the same number of moles in the same volume, so their

concentrations are the same.

- (d) According to kinetic-molecular theory, the average kinetic energy of a gas depends only on the temperature. Both gases are at the same temperature, so they have the same average kinetic energy.
- (e) H₂O will deviate most from ideal behavior. Ideal behavior for gas molecules assumes that there will be no intermolecular interactions.

H₂O is polar, and O₂ is not. H₂O undergoes hydrogen bonding, while O₂ does not. So H₂O has stronger intermolecular interactions, which will cause it to deviate more from ideal behavior.

8. (a) $745 \text{ mmHg} - 16.5 \text{ mmHg} = 729 \text{ mmHg}$

$$\frac{729 \text{ mmHg}}{760 \text{ mmHg}} = 0.959 \text{ atm}$$

- (b) To determine the mass of the butane, subtract the mass of the lighter after the butane was released from the mass of the lighter before the butane was released.

$$20.432 \text{ g} - 20.296 \text{ g} = 0.136 \text{ g}$$

To determine the moles of butane, use the Ideal Gas Law, making any necessary conversions first.

$$PV = nRT$$

$$(0.959 \text{ atm})(0.06840 \text{ L}) = n(0.0821 \text{ atm}\cdot\text{L/mol}\cdot\text{K})(292 \text{ K})$$

$$n = 2.74 \times 10^{-3} \text{ mol}$$

Molar mass is defined as grams per mole, so

$$0.136 \text{ g} / 2.74 \times 10^{-3} \text{ mol} = 49.6 \text{ g/mol}$$

(c) Actual molar mass of butane:

$$(12.00 \text{ g/mol} \times 4) + (1.01 \text{ g/mol} \times 10) = 58.08 \text{ g/mol}$$

Percent error is:

$$\frac{|\text{Actual value} - \text{experimental value}|}{\text{Actual value}} \times 100\%$$

So:

$$\frac{|58.08 - 49.6|}{58.08} \times 100\% = 14.5\% \text{ error}$$

- (d) (i) If the lighter is not sufficiently dried, then the mass of the butane calculated will be artificially low. That means the numerator in the molar mass calculation will be too low, which would lead to an experimental molar mass that is too low. This is consistent with the student's error.
- (ii) If the temperature of the butane is higher than the water temperature, the calculated moles of butane will be artificially high. This means the denominator in the molar mass calculation will be too high, which would lead to an experimental molar mass that is too low. This is consistent with the student's error.
- (iii) If some butane gas escaped without going into the graduated cylinder, the volume of butane gas collected will be artificially low. That will make the calculation for moles of butane too low, which in turns means the denominator of the molar mass calculation will be too low. This would lead to an experimental molar mass that

is too high. This is NOT consistent with the student's error.

Chapter 5

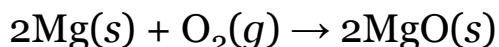
Big Idea #3: Chemical Reactions, Energy Changes, and Redox Reactions

Changes in matter involve the rearrangement and/or reorganization of atoms and/or the transfer of electrons.

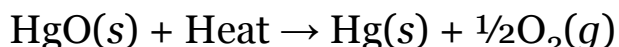
TYPES OF REACTIONS

Reactions may be classified into several categories.

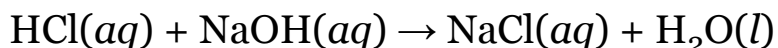
1. **Synthesis Reactions:** When elements or simple compounds are combined to form a single, more complex compound.



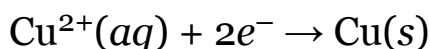
2. **Decomposition:** The opposite of a synthesis. A reaction where a single compound is split into two or more elements or simple compounds, usually in the presence of heat.



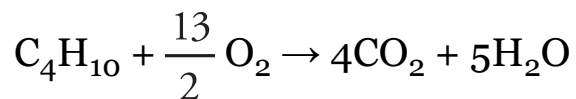
3. **Acid-Base Reaction:** A reaction when an acid (i.e., H^+) reacts with a base (i.e., OH^-) to form water and a salt.



4. **Oxidation-Reduction (Redox) Reaction:** A reaction that results in the change of the oxidation states of some participating species.

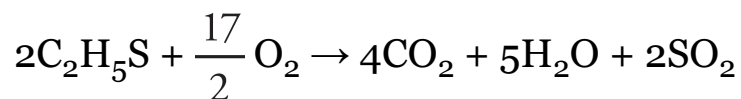


5. **Hydrocarbon Combustion:** When a covalent substance containing carbon and hydrogen (and sometimes oxygen) is ignited, it reacts with the oxygen in the atmosphere. The products of a hydrocarbon combustion are always CO_2 and H_2O . The combustion of butane, a fuel found in many lighters, is an example of a combustion reaction:



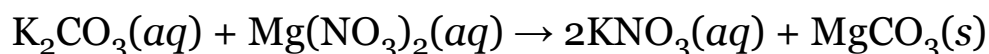
If elements other than carbon and hydrogen are present in the substance being combusted, they, too, often combine with

oxygen to form various gases.

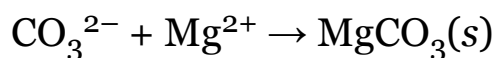


It is not always possible to predict the exact formula of the non-carbon or hydrogen containing compounds that are created, but carbon will always lead to CO_2 and hydrogen, to H_2O .

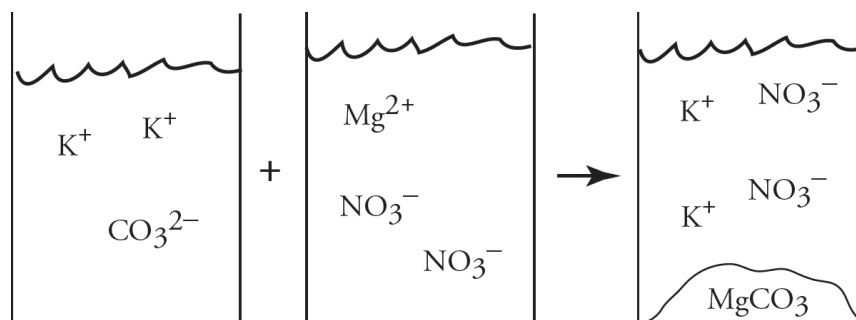
6. Precipitation: When two aqueous solutions mix, sometimes a new cation/anion pairing can create an insoluble salt. This type of reaction is called a precipitation reaction. When potassium carbonate and magnesium nitrate mix, a precipitate of magnesium carbonate will form as follows:



That can also be written as a net ionic equation. In solution, the potassium and nitrate ions do not actually take part in the reaction. They start out as free ions and end up as free ions. We call those ions **spectator ions**. The only thing that is changing is that carbonate and magnesium ions are bonding to form magnesium carbonate. The net ionic equation is:



Finally, this can be represented using particulate diagrams, as shown below:

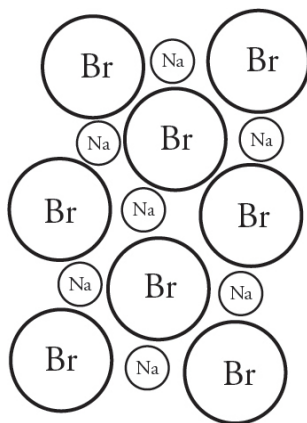


You may have heard that you need to memorize solubility rules—that is, what ions will form insoluble precipitates when you combine them. You DO NOT need to memorize most of these; for the most part, the AP Exam will provide you with them as needed. The only ones you need to know are:

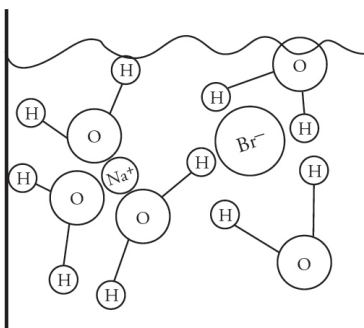
1. Compounds with an alkali metal cation (Na^+ , Li^+ , K^+ , etc.) or an ammonium cation (NH_4^+) are always soluble.
2. Compounds with a nitrate (NO_3^-) anion are always soluble.

That's it! You should know how to interpret any solubility rules that you are given on the test, but you need not memorize anything beyond those two.

Ionic substances that dissolve in water do so because of the attraction of the ions to the dipoles of the water molecules exceeds the attraction of the ions to each other. Let's take NaBr for example:



When it dissolves in water, the positive Na^+ cations are attracted to the negative (oxygen) ends of the water molecules. The negative Br^- anions are attracted to the positive (hydrogen) ends of the water molecules. Thus, when the substance dissolves it looks like this on the particulate level:



You should be familiar with the following polyatomic ions and their charges.

Hydroxide	OH^-
Nitrate	NO_3^-
Acetate	$\text{C}_2\text{H}_3\text{O}_2^-$
Cyanide	CN^-
Permanganate	MnO_4^-
Carbonate	CO_3^{2-}
Sulfate	SO_4^{2-}
Dichromate	$\text{Cr}_2\text{O}_7^{2-}$
Phosphate	PO_4^{3-}
Ammonium	NH_4^+



More Great Books

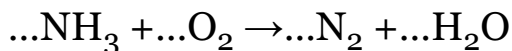
Check out another book from The Princeton Review about chemistry—*High School Chemistry Unlocked* for more content review and practice.

CHEMICAL EQUATIONS

Balancing Chemical Equations

Normally, balancing a chemical equation is a trial-and-error process. You start with the most complicated-looking compound in the equation and work from there. There is, however, an old Princeton Review SAT trick that you may want to try if you see a balancing equation question on the multiple-choice section. The trick is called **backsolving**.

It works like this: To make a balancing equation question work in a multiple-choice format, one of the answer choices is the correct coefficient for one of the species in the reaction. So instead of starting blind in the trial-and-error process, you can insert the answer choices one by one to see which one works. You probably won't have to try all four, and if you start in the middle and the number doesn't work, it might be obviously too small or large, eliminating other choices before you have to try them. Let's try it.



1. If the equation above were balanced with lowest whole number coefficients, the coefficient for NH_3 would be

- (A) 1
- (B) 2
- (C) 3
- (D) 4

Here's How to Crack It!

Start at (C) because it's a middle number. If there are 3 NH_3 's, then there can't be a whole number coefficient for N_2 , so (C) is wrong, and so is the other odd number answer, (A).

Try (D).

If there are 4 NH_3 's, then there must be 2 N_2 's and 6 H_2O 's.

If there are 6 H_2O 's, then there must be 3 O_2 's, and the equation is balanced with lowest whole number coefficients.

A Note About Backsolving

Backsolving is more efficient than the methods that you're accustomed to. If you use the answer choices that you're given, you streamline the trial-and-error process and allow yourself to use POE as you work on the problem.

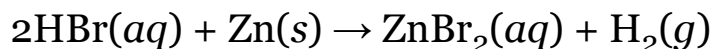
Chemical Equations and Calculations

Because of the math involved, stoichiometry like this will most likely appear only as a free-response question. In those instances, expect the following: You will be given a balanced chemical equation and told that you have some number of grams (or liters of gas, or molar concentration, and so on) of reactant. Then you will be asked what number of grams (or liters of gas, or molar concentration, and so on) of products are generated.

In these cases, follow this simple series of steps.

1. Convert whatever quantity you are given into moles.
2. If you are given information about two reactants, you may have to use the equation coefficients to determine which one is the limiting reagent. Remember, the limiting reagent is not necessarily the reactant that you have the least of; it is the reactant that runs out first.
3. Use the balanced equation to determine how many moles of the desired product are generated.
4. Convert moles of product to the desired unit.

Let's try one.



2. A piece of solid zinc weighing 98 grams was added to a solution containing 324 grams of HBr. What is the volume of H_2 produced at standard temperature and pressure if the reaction above runs to completion?

Here's How to Crack It!

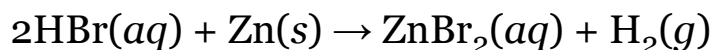
1. Convert whatever quantity you are given into moles.

$$\text{Moles of Zn} = \frac{\text{grams}}{\text{molar mass}} = \frac{(98 \text{ g})}{(65.4 \text{ g/mol})} = 1.5 \text{ mol}$$

$$\text{Moles of HBr} = \frac{\text{grams}}{\text{molar mass}} = \frac{(324 \text{ g})}{(80.9 \text{ g/mol})} = 4.0 \text{ mol}$$

2. Use the balanced equation to find the limiting reagent. From the balanced equation, 2 moles of HBr are used for every mole of Zn that reacts, so when 1.5 moles of Zn react, 3 moles of HBr are consumed, and there will be HBr left over when all of the Zn is gone. That makes Zn the limiting reagent.
3. Use the balanced equation to determine how many moles of the desired product are generated.
1 mole of H₂ is produced for every mole of Zn consumed, so if 1.5 moles of Zn are consumed, then 1.5 moles of H₂ are produced.
4. Convert moles of product to the desired unit.
The H₂ gas is at STP, so you can convert directly from moles to volume. Volume of H₂ = (moles)(22.4 L/mol) = (1.5 mol)(22.4 L/mol) = 33.6 L $\approx$ 34 L

Let's try another one using the same reaction.



3. A piece of solid zinc weighing 13.1 grams was placed in a container. A 0.10-molar solution of HBr was slowly added to the container until the zinc was completely dissolved. What was the volume of HBr solution required to completely dissolve the solid zinc?

Here's How to Crack It!

1. Convert whatever quantity you are given into moles.

$$\text{Moles of Zn} = \frac{\text{grams}}{\text{MW}} = \frac{(13.1 \text{ g})}{(65.4 \text{ g/mol})} = 0.200 \text{ mol}$$

2. Use the balanced equation to find the limiting reagent.
3. Use the balanced equation to determine how many moles of the desired product are generated.

In this case, use the balanced reaction to find out how much of one reactant is required to consume the other reactant.

You can see from the balanced equation that it takes 2 moles of HBr to react completely with 1 mole of Zn, so it will take 0.400 moles of HBr to react completely with 0.200 moles of Zn.

4. Convert moles of product to the desired unit. Moles of HBr = (molarity)(volume)

$$\text{Volume of HBr} = \frac{\text{moles}}{\text{molarity}} = \frac{(0.400 \text{ mol})}{(0.10 \text{ mol/L})} = 4.0 \text{ L}$$

When you perform calculations, always include units. Including units in your calculations will help you (and the person scoring your test) keep track of what you are doing. While using the proper units in an answer is not typically worth any points in and of itself (with a few exceptions), AP graders may take off points if your units are missing or incorrect.

Combustion Analysis

When a hydrocarbon is combusted, producing carbon dioxide and water, the formula of that hydrocarbon can often be determined using a little math. One of the most important concepts in chemistry is the **Law of Conservation of Mass**. This states that matter can

be neither created nor destroyed during a chemical reaction. Thus, when a hydrocarbon is combusted, all of the carbon in that hydrocarbon will end up in CO_2 , and all of the hydrogen will end up in H_2O . With that in mind, you can often determine the empirical formula of a hydrocarbon by determining the mass of the products it forms.

4. A compound containing hydrogen, carbon, and oxygen is combusted. 44.0 g of CO_2 and 27.0 g of H_2O are produced. Which of the following is a possible empirical formula for that compound?

- (A) CH_4O
- (B) $\text{C}_2\text{H}_6\text{O}$
- (C) $\text{C}_2\text{H}_3\text{O}_2$
- (D) $\text{C}_3\text{H}_5\text{O}_2$

Here's How to Crack It!

If 44.0 grams of CO_2 are created, that is 1.00 mole of CO_2 , and thus, 1.00 mole of C as well (as there is one mole of carbon in each mole of CO_2). Through conservation of mass, this means there was 1.00 mole of C in the original compound.

27.0 g of H_2O means 1.5 moles of H_2O were created. Because there are two hydrogen atoms in every H_2O , that means there were 3.0 moles of hydrogen in the water which was created, and thus, 3.00 moles of H in the original compound. The C:H ratio in the original compound must be 1:3, which is consistent with (B).

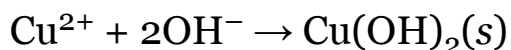
GRAVIMETRIC ANALYSIS

A common way to use precipitation reactions in order to make quantitative determinations about the identity of an unknown sample is through gravimetric analysis. In this process, a soluble sample of an unknown compound is mixed with another solution that will cause one of the ions from the unknown sample to fully precipitate out. If the identity of the precipitate is known, stoichiometric conversions will reveal the mass of the ion in the unknown, which allows for a mass percent calculation.

Due to the large amount of math involved, it's easiest to demonstrate.

Example:

A 4.33 g sample of an unknown alkali hydroxide compound is dissolved completely in water. A sufficient solution of copper (II) nitrate is added to the hydroxide solution such that it will fully precipitate copper (II) hydroxide via the following reaction:



After the precipitate is filtered and dried, its mass is found to be 3.81 g. Is the original alkali hydroxide sample most likely LiOH, NaOH, or KOH?

First, you convert the mass of the precipitate to moles:

$$3.81\text{g Cu}(\text{OH})_2 \times \frac{1 \text{ mol Cu}(\text{OH})_2}{97.55\text{g Cu}(\text{OH})_2} = 0.0391 \text{ mol Cu}(\text{OH})_2$$

Next, you need to determine the moles of hydroxide in the precipitate and convert that to grams.

$$0.0391 \text{ mol Cu}(\text{OH})_2 \times \frac{2 \text{ mol OH}^{-}}{1 \text{ mol Cu}(\text{OH})_2} \times \frac{17.02 \text{ g OH}^{-}}{1 \text{ mol OH}^{-}} = 1.33 \text{ g OH}^{-}$$

If there are 1.33 g of OH^- present in the precipitate, then there were also 1.33 g of OH^- in the original sample. The final calculation is to determine the mass percent of the OH^- in the original sample:

$$\frac{1.33 \text{ g OH}^-}{4.33 \text{ g sample}} \times 100\% = 30.7\%$$

To determine the identity of the sample, you have to know the mass percent of hydroxide in all of the possible compounds. To do that, you divide the mass of one mole of hydroxide by the molar mass of the compound.

$$\text{LiOH: } \frac{17.02 \text{ g}}{23.96 \text{ g}} \times 100\% = 71.0\%$$

$$\text{NaOH: } \frac{17.02 \text{ g}}{40.02 \text{ g}} \times 100\% = 42.5\%$$

$$\text{KOH: } \frac{17.02 \text{ g}}{56.12 \text{ g}} \times 100\% = 30.3\%$$

The hydroxide mass percent in KOH is closest to the mass percent determined via gravimetric analysis, so that is the most likely identity of the original unknown.

ENTHALPY

Enthalpy Change, ΔH

The enthalpy of a substance is a measure of the energy that is released or absorbed by the substance when bonds are broken and formed during a reaction.

The Basic Rules of Enthalpy

When bonds are **formed**, energy is **released**.
When bonds are **broken**, energy is **absorbed**.

The change in enthalpy, ΔH , that takes place over the course of a reaction can be calculated by subtracting the enthalpy of the reactants from the enthalpy of the products.

Enthalpy Change

$$\Delta H = H_{\text{products}} - H_{\text{reactants}}$$

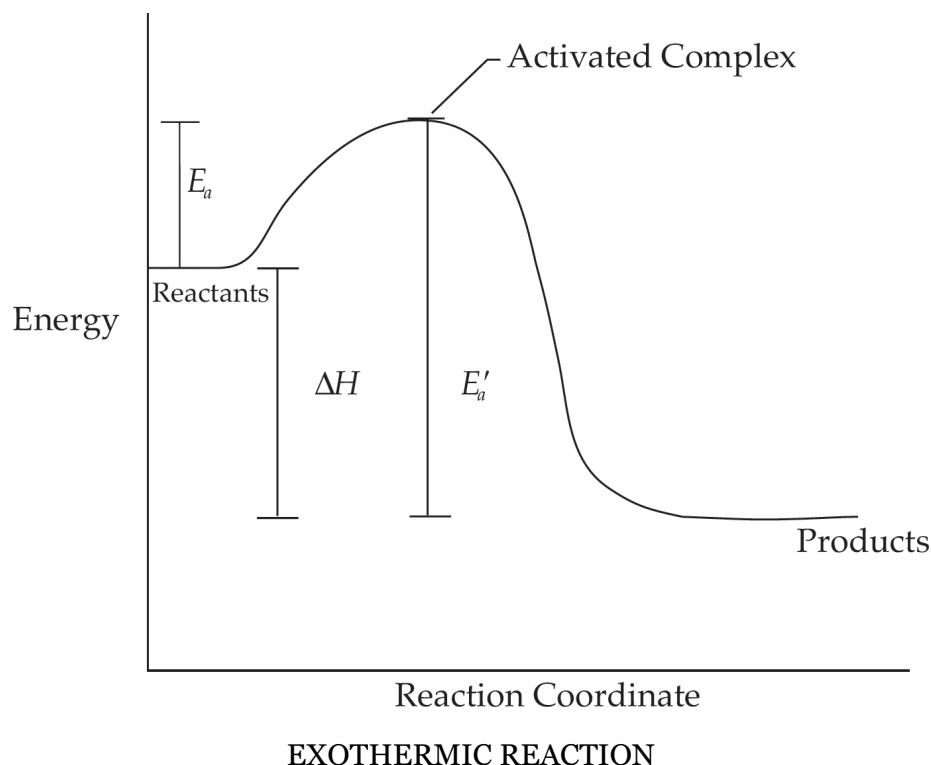
If more energy is released when the bonds in the products form than is necessary to break the bonds in the reactants, the overall reaction releases energy into the surroundings and the reaction is called **exothermic**. Exothermic reactions have a negative enthalpy change.

If more energy is required to break the bonds in the reactants than is released when the bonds in the products form, the overall reaction requires energy from the surroundings and is called **endothermic**. Endothermic reactions have a positive enthalpy change.

All substances are the most stable in their lowest energy state. This means that, in general, exothermic processes are more likely to be favorable than endothermic processes.

ENERGY DIAGRAMS

Exothermic and Endothermic Reactions

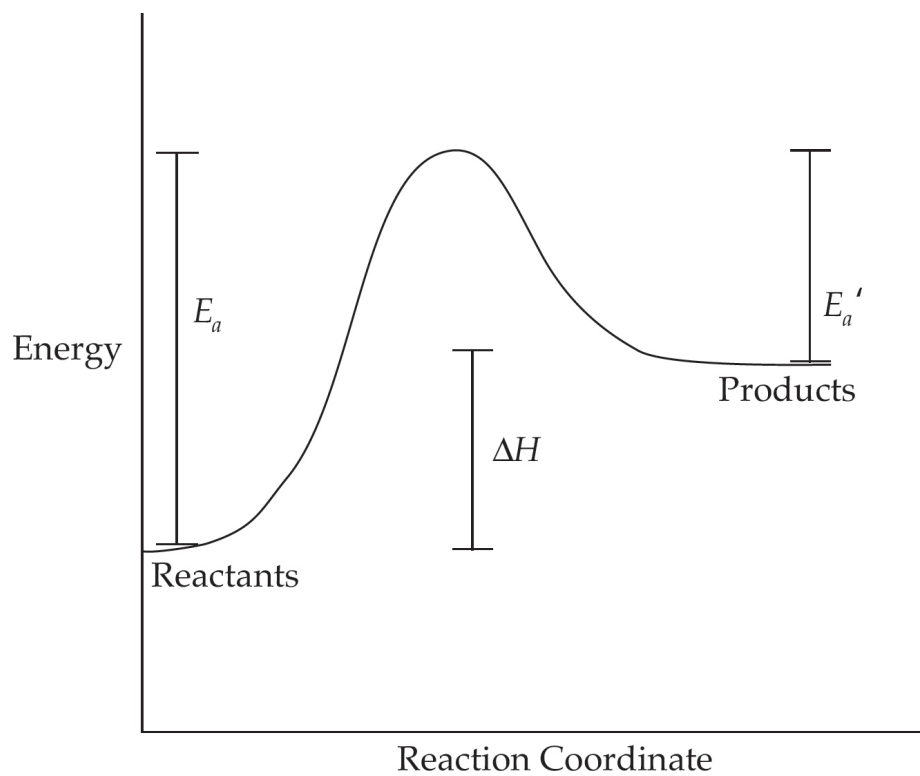


The diagram above shows the energy change that takes place during an exothermic reaction. The reactants start with a certain amount of energy (read the graph from left to right). For the reaction to proceed, the reactants must have enough energy to reach the transition state, where they are part of an activated complex. This is the highest point on the graph above. The amount of energy needed to reach this point is called the **activation energy**, E_a . At this point, all reactant bonds have been broken, but no product bonds have been formed, so this is the point in the reaction with the highest energy and lowest stability. The energy needed for the reverse reaction is shown as line E'_a .

Moving to the right past the activated complex, product bonds start to form, and we eventually reach the energy level of the products.

This diagram represents an exothermic reaction, so the products are at a lower energy level than the reactants and ΔH is negative.

The diagram below shows an endothermic reaction.

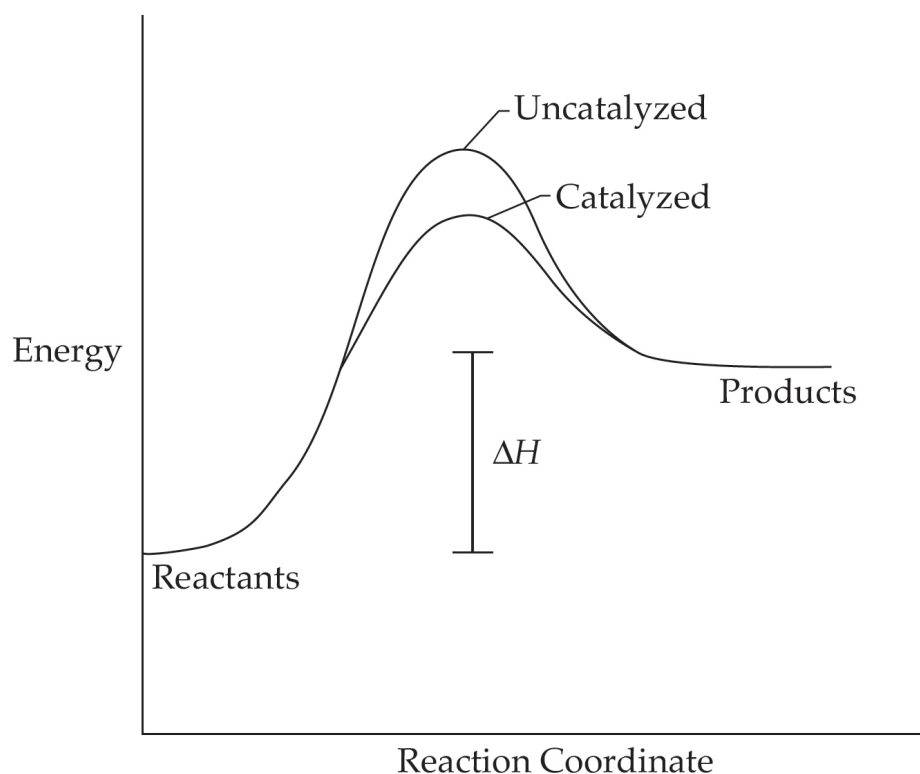


ENDOTHERMIC REACTION

In this diagram, the energy of the products is greater than the energy of the reactants, so ΔH is positive.

Reaction diagrams can be read in both directions, so the reverse reaction for an exothermic reaction is endothermic and vice versa.

CATALYSTS AND ENERGY DIAGRAMS



A catalyst speeds up a reaction by providing the reactants with an alternate pathway that has a lower activation energy, as shown in the diagram above.

Notice that the only difference between the catalyzed reaction and the uncatalyzed reaction is that the energy of the activated complex is lower for the catalyzed reaction. A catalyst lowers the activation energy, but it has no effect on the energy of the reactants, the energy of the products, or ΔH for the reaction.

Also note that a catalyst lowers the activation energy for both the forward and the reverse reaction, so it has no effect on the equilibrium conditions.

OXIDATION STATES

In many chemical reactions, electrons are transferred between the reactants. In order to determine which reactants are gaining electrons and which are losing electrons, we use something called an

oxidation state. There are several rules when it comes to assigning oxidation states.

- 1) Any neutral atom that is not bonded to atoms of any other element has an oxidation state of zero. Examples: In Cu(s) , the oxidation state on copper is zero. In $\text{O}_2(g)$, the oxidation state on both oxygen atoms is zero.
- 2) Any ion has an oxidation state equal to the charge on that ion. This includes ions bonded to other ions in any kind of ionic compound. Examples: In $\text{Zn}^{2+}(aq)$ zinc has an oxidation state of +2. In FeCl_3 , iron is +3 and chlorine is -1.
- 3) In most compounds, oxygen is -2. An exception worth knowing is that in hydrogen peroxide (H_2O_2) oxygen is -1. Example: In $\text{C}_6\text{H}_{12}\text{O}_6$, oxygen is -2.
- 4) When bonded to a nonmetal, hydrogen is +1. When bonded to a metal, hydrogen is -1, Examples: In CH_4 , hydrogen is +1. In NaH , hydrogen is -1.
- 5) In the absence of oxygen, the most electronegative element in a compound will take on an oxidation state equal to its most common charge. Examples: In CF_4 , fluorine is -1. In CS_2 , sulfur is -2.
- 6) The combined oxidation states on all elements in a neutral compound must add up to zero. The combined oxidation states on all elements in a polyatomic ion must add up to the charge on that ion.

Rule 6 is only applied for covalent compounds to determine the oxidation state on an element which does not fall under the first five rules. Not that many nonmetals with low electronegativity values (such as carbon, nitrogen, sulfur, and phosphorus) have frequently varying oxidation states.

Example 1: Determine the oxidation state on every element in H_2SO_4 .

We know that H is +1 and O is -2, and that the total charge on the compound must be 0. Thus:

$$+1(2) + S + -2(4) = 0.$$

Solve for S, and you will find $S = +6$.

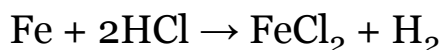
Example 2: Determine the oxidation state on every element in the phosphate ion, PO_4^{3-} .

$$P + -2(4) = -3$$

$$P = +5$$

OXIDATION-REDUCTION REACTIONS

In an oxidation-reduction (or redox, for short) reaction, electrons are exchanged by the reactants, and the oxidation states of some of the reactants are changed over the course of the reaction. Look at the following reaction:



The oxidation state of Fe changes from 0 to +2.

The oxidation state of H changes from +1 to 0.

- *When an atom gains electrons, its oxidation number decreases, and it is said to have been reduced.*

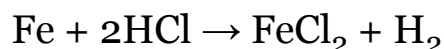
In the reaction above, H was reduced.

- *When an atom loses electrons, its oxidation number increases, and it is said to have been oxidized.*

In the reaction above, Fe was oxidized.

Oxidation and reduction go hand in hand. If one atom is losing electrons, another atom must be gaining them.

An oxidation-reduction reaction can be written as two **half-reactions**: one for the reduction and one for the oxidation. For example, the reaction



can be written as



Here's a mnemonic device that might be useful.

LEO the lion says GER

LEO: you **L**ose **E**lectrons in **O**xidation

GER: you **G**ain **E**lectrons in **R**eduction

Or, if lions don't appeal to you, another way to memorize this concept is **OIL RIG**:

Oxidation **I**s **L**oss

Reduction **I**s **G**ain

Redox Titrations

A titration involves the slow addition of a solution at a known concentration to another solution of unknown concentration in order to determine the concentration of the unknown solution. To determine the endpoint of a titration, a color change is often used. Titrations are frequently used in acid-base reactions (more on that in

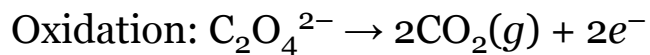
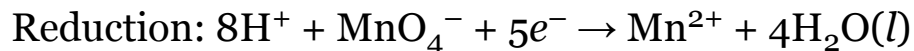
Chapter 8), but redox reactions can also be used in titration situations.

Potassium permanganate, KMnO_4 , is frequently used in redox titrations. The manganese ion has an oxidation state of +7 in the permanganate ion (MnO_4^-), and a solution containing permanganate ions is a deep purple color. The manganese in MnO_4^- reduces to Mn^{2+} (thus changing its oxidation state to +2) when mixed with compounds that can be oxidized. Mn^{2+} is a colorless ion.

Frequently, when potassium permanganate is titrated into a colorless solution that contains a compound that can be oxidized, the end of the titration is marked by the solution turning pink. Initially, the permanganate ions take electrons from the oxidized compound and reduce to Mn^{2+} . However, once the compound that is being oxidized runs out, there are no electrons left for the MnO_4^- to take, and thus any extra permanganate ion that is added remains unreduced and retains its purple color, which when diluted sufficiently appears pink.

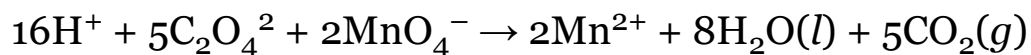
Example:

A 50.0 mL solution of sodium oxalate, $\text{Na}_2\text{C}_2\text{O}_4$ is poured into an Erlenmeyer flask. An acidified solution of 0.135 *M* potassium permanganate is titrated into the flask while the solution is swirled on a stir station and heated. The solution in the flask is originally colorless, but turns pink after 14.56 mL of potassium permanganate is added. Given the following half-reactions, what is the concentration of the oxalate solution?



The first step to solving this problem is to balance the half-reactions. In this case, the oxidation half-reaction must be multiplied by 5 in

order to make the electrons balance with those in the reduction half-reaction, and the reduction half-reaction must be multiplied by 2. Thus, the full reaction (which omits the sodium and potassium spectator atoms) is:



So, for every mole of permanganate that reacts, five moles of oxalate are required. The number of moles of permanganate can be easily determined:

$$M = \frac{n}{V} 0.135 \text{ M MnO}_4^- = \frac{n}{0.01456 \text{ L MnO}_4^-} n = 0.00197 \text{ MnO}_4^-$$

That can then be converted to moles of oxalate:

$$0.00197 \text{ MnO}_4^- \times \frac{5 \text{ mol C}_2\text{O}_4^{2-}}{2 \text{ mol MnO}_4^-} = 0.00493 \text{ mol C}_2\text{O}_4^{2-}$$

Finally, the concentration of the oxalate solution can be determined:

$$\frac{0.00493 \text{ mol C}_2\text{O}_4^{2-}}{0.0500 \text{ L C}_2\text{O}_4^{2-}} = 0.0986 \text{ M} = [\text{C}_2\text{O}_4^{2-}] = [\text{Na}_2\text{C}_2\text{O}_4^{2-}]$$

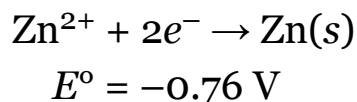
You will not be expected to memorize any of the various color changes that can occur throughout a redox titration. However, you should understand the concept of using a color change to determine an endpoint if you are provided with the colors of the various compounds that are present during a titration.

Reduction Potentials

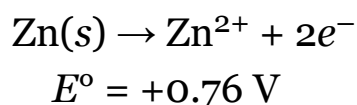
Every half-reaction has an electric potential, or voltage, associated with it. You will be given the necessary values for the standard reduction potential of half-reactions for any question in which they are required. Potentials are always given as reduction half-reactions,

but you can read them in reverse and flip the sign on the voltage to get oxidation potentials.

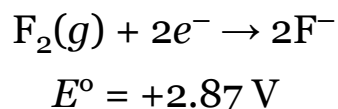
Look at the reduction potential for Zn^{2+} .



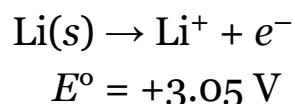
Read the reduction half-reaction in reverse and change the sign on the voltage to get the oxidation potential for Zn.



The larger the potential for a half-reaction, the more likely it is to occur.



$\text{F}_2(g)$ has a very large reduction potential, so it is likely to gain electrons and be reduced.



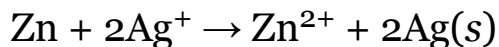
$\text{Li}(s)$ has a very large oxidation potential, making it very likely to lose electrons and be oxidized.

You can calculate the potential of a redox reaction if you know the potentials for the two half-reactions that constitute it. There are two important things to remember when calculating the potential of a redox reaction.

- Add the potential for the oxidation half-reaction to the potential for the reduction half-reaction.

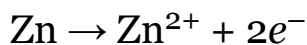
- Never multiply the potential for a half-reaction by a coefficient.

Let's look at the following reaction:



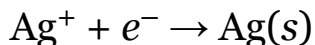
The two half-reactions are:

Oxidation:



$$E^\circ = +0.76 \text{ V}$$

Reduction:



$$E^\circ = +0.80 \text{ V}$$

$$E = E_{\text{oxidation}} + E_{\text{reduction}}$$

$$E = 0.76 \text{ V} + 0.80 \text{ V} = 1.56 \text{ V}$$

Notice that we ignored that silver has a coefficient of 2 in the balanced equation.

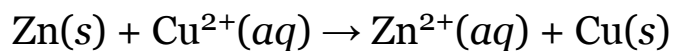
The relative reduction strengths of two different metals can also be determined qualitatively. In the above reaction, if Zn(s) were placed in a solution containing Ag⁺ ions, the silver ions have a high enough reduction potential that they would take electrons from the zinc and start forming solid silver, which would precipitate out on the surface of the zinc.

However, if Ag(s) were placed in a solution containing Zn²⁺ ions, zinc does not have a high enough reduction potential to take electrons from silver and so no reaction would occur. So, when a solid metal is placed into a metallic solution and a new solid starts to form, the reduction potential of the metal in solution is greater than that of the solid metal. If no solid forms, the reduction potential of the solid metal is higher.

GALVANIC CELLS

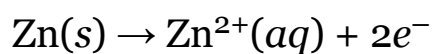
In a **galvanic cell** (also called a voltaic cell), a favored redox reaction is used to generate a flow of current.

Look at the following favored redox reaction:



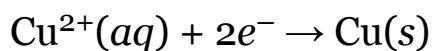
$$E^{\circ} = 1.10 \text{ V}$$

Oxidation:



$$E^{\circ} = 0.76 \text{ V}$$

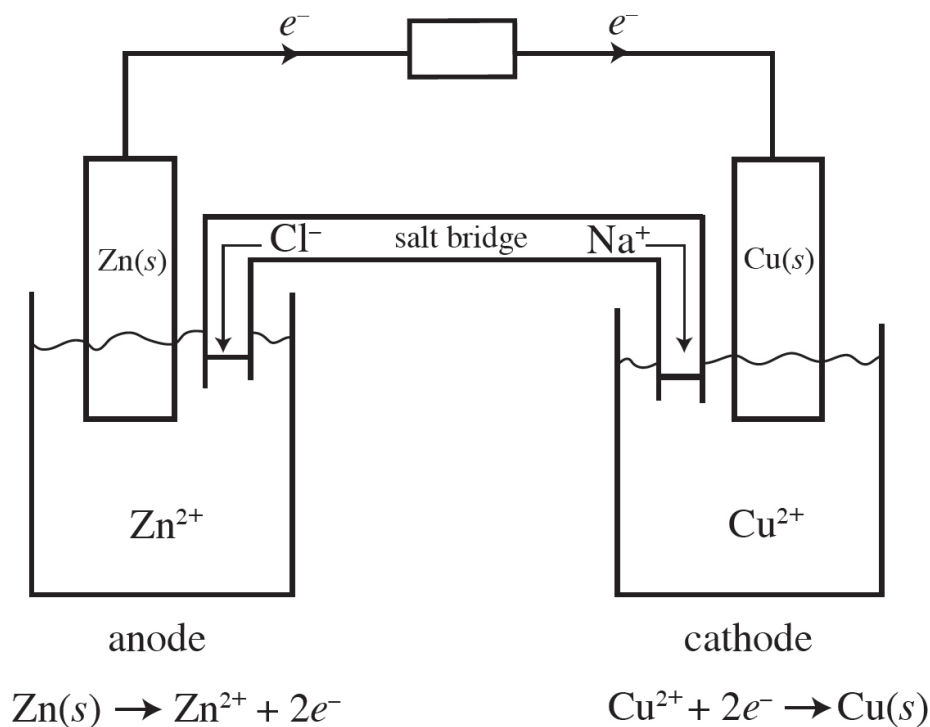
Reduction:



$$E^{\circ} = 0.34 \text{ V}$$

Going Platinum

Because it is unreactive with most solutions, platinum is often used as a conductor in an electrolytic cell.



In a galvanic cell, the two half-reactions take place in separate chambers, and the electrons that are released by the oxidation reaction pass through a wire to the chamber where they are consumed in the reduction reaction. That's how the current is created. **Current** is defined as the flow of electrons from one place to another.

In any electric cell (either a galvanic cell or an electrolytic cell, which we'll discuss a little later) oxidation takes place at the electrode called the **anode**. Reduction takes place at the electrode called the **cathode**.

There's a mnemonic device to remember that.

AN OX
RED CAT

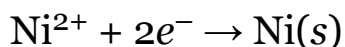
The salt bridge maintains the electrical neutrality in the cell. At the cathode, where reduction occurs and the solution is becoming less positively charged, the positive cations from the salt bridge solution flow into the half-cell. At the anode, where oxidation occurs and the solution is becoming more positively charged, the negative anions from the salt bridge solution flow into the half-cell. If the salt bridge were to be removed, the solutions in the half-cells would become electrically imbalanced, and the voltage of the cell would drop to 0. In the previous diagram, the salt bridge is filled with aqueous NaCl. As the cell functions, the Na^+ cations flow to the cathode and the Cl^- anions flow to the anode.

Under standard conditions (when all concentrations are 1 *M*), the voltage of the cell is the same as the total voltage of the redox reaction. To determine what would happen to the cell potential under nonstandard conditions, you must first learn more about equilibrium, which will be covered in Chapter 8.

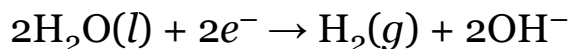
ELECTROLYTIC CELLS

In an electrolytic cell, an outside source of voltage is used to force an unfavored redox reaction to take place. Most electrolytic cells occur in aqueous solutions, which are created when a chemical dissolves in water; either the ions or the water molecules can be reduced or oxidized.

Let's look at a solution of nickel (II) chloride as an example. To determine which substance is reduced, you must compare the reduction potential of the cation with that of water. The half-reaction with the more positive value is the one that will occur.

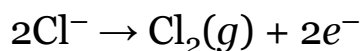


$$E^\circ = -0.25 \text{ V}$$

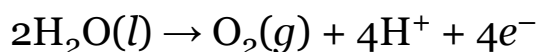


$$E^\circ = -0.80 \text{ V}$$

In this case, the Ni^{2+} reduction will occur. For the oxidation, the oxidation potential of the anion versus that of water must be examined. As with before, the half-reaction with the most positive value is the one that will occur. Remember that potentials are always given as reduction potentials, so you must flip the sign when you flip the equation to an oxidation.

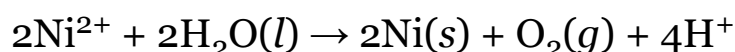


$$E^\circ = -1.36 \text{ V}$$



$$E^\circ = -1.23 \text{ V}$$

So, in this case, the water itself would be reduced. When the equations are balanced for electron transfer, the net ionic equation looks like this:



$$E = -0.25 \text{ V} + -1.24 \text{ V} = -1.49 \text{ V}$$

The anode and cathode in an electrolytic reaction are usually just metal bars that conduct current and do not take part in the reaction. In the above reaction, solid nickel is being created at the cathode, and oxygen gas is being evolved at the anode. The sign of your total cell potential (E) for an electrolytic reaction is always negative.

Occasionally, a current will either be run through a molten compound or pure water. In this case, you do not need to determine which redox reactions are taking place, as you will only have one choice for each.

The AN OX/RED CAT rule applies to the electrolytic cell in the same way that it applies to the galvanic cell.

Electroplating

Electrolytic cells are used for electroplating. You may see a question on the test that gives you an electrical current and asks you how much metal “plates out.”

There are roughly four steps for figuring out electrolysis problems.

1. If you know the current and the time, you can calculate the charge in coulombs.

Current

$$I = \frac{q}{t}$$

I = current (amperes, A)

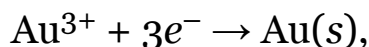
q = charge (coulombs, C)

t = time (seconds, s)

2. Once you know the charge in coulombs, you know how many electrons were involved in the reaction.

$$\text{Moles of electrons} = \frac{\text{coulombs}}{96,500 \text{ coulombs/mol}}$$

3. When you know the number of moles of electrons and you know the half-reaction for the metal, you can find out how many moles of metal plated out. For example, from this half-reaction for gold,



you know that for every 3 moles of electrons consumed, you get 1 mole of gold.

4. Once you know the number of moles of metal, you can use what you know from stoichiometry to calculate the number of grams of metal.

For instance, if a current of 2.50 A is run through a solution of iron (III) chloride for 15 minutes, it would cause the following mass of iron to plate out:

$$15 \text{ minutes} \times \frac{60 \text{ seconds}}{1 \text{ minute}} \times \frac{2.50 \text{ C}}{1.0 \text{ second}} \times \frac{1 \text{ mol } e^{-}}{96,500 \text{ C}} \times \frac{1 \text{ mol Fe}}{3 \text{ mol } e^{-}} \times \frac{55.85 \text{ g}}{1 \text{ mol Fe}} = 0.434 \text{ g Fe}$$

It is particularly important to keep track of your units in an electroplating problem; there are many different conversions before you come up with your final answer. In general, as long as all of your conversions are set up correctly your final answer will have the correct units.

CHAPTER 5 QUESTIONS

[Click here](#) to download and print the PDF version of this exercise.

Multiple-Choice Questions

Use the following solubility rules to answer questions 1-4.

Salts containing halide anions are soluble except for those containing Ag^+ , Pb^{2+} , and Hg_2^{2+} . Salts containing carbonate anions are insoluble except for those containing alkali metals or ammonium.

1. If solutions of iron (III) nitrate and sodium carbonate are mixed, what would be the formula of the precipitate?

(A) Fe_3CO_3
(B) $\text{Fe}_2(\text{CO}_3)_3$
(C) NaNO_3
(D) No precipitate would form.
2. If solutions containing equal amounts of AgNO_3 and KCl are mixed, what is the identity of the spectator ions?

(A) Ag^+ , NO_3^- , K^+ , and Cl^-
(B) Ag^+ and Cl^-
(C) K^+ and Ag^+
(D) K^+ and NO_3^-

3. If equimolar solutions of $\text{Pb}(\text{NO}_3)_2$ and NaCl are mixed, which ion will not be present in significant amounts in the resulting solution after equilibrium is established?
- (A) Pb^{2+}
(B) NO_3^-
(C) Na^+
(D) Cl^-
4. Choose the correct net ionic equation representing the reaction that occurs when solutions of potassium carbonate and copper (I) chloride are mixed.
- (A) $\text{K}_2\text{CO}_3(\text{aq}) + 2\text{CuCl}(\text{aq}) \rightarrow 2\text{KCl}(\text{aq}) + \text{Cu}_2\text{CO}_3(\text{s})$
(B) $\text{K}_2\text{CO}_3(\text{aq}) + 2\text{CuCl}(\text{aq}) \rightarrow 2\text{KCl}(\text{s}) + \text{Cu}_2\text{CO}_3(\text{aq})$
(C) $\text{CO}_3^{2-} + 2\text{Cu}^+ \rightarrow \text{Cu}_2\text{CO}_3(\text{s})$
(D) $\text{CO}_3^{2-} + \text{Cu}^{2+} \rightarrow \text{CuCO}_3(\text{s})$
5. A strip of metal X is placed into a solution containing Y^{2+} ions and no reaction occurs. When metal X is placed in a separate solution containing Z^{2+} ions, metal Z starts to form on the strip. Which of the following choices organizes the reduction potentials for metals X, Y, and Z from greatest to least?
- (A) $\text{X} > \text{Y} > \text{Z}$
(B) $\text{Y} > \text{Z} > \text{X}$
(C) $\text{Z} > \text{X} > \text{Y}$
(D) $\text{Y} > \text{X} > \text{Z}$
6. In which of the following compounds is the oxidation number of chromium the greatest?



7. For an endothermic reaction, which of the following is true regarding the energy level of the activated complex?

(A) It is above the energy level of the reactants, but below the energy level of the products.

(B) It is below the energy level of the reactants, but above energy level of the products.

(C) It is above the energy level of both the products and reactants.

(D) It is below the energy level of both the products and reactants.

8. What is the mass of oxygen in 148 grams of calcium hydroxide (Ca(OH)_2)?

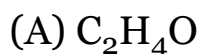
(A) 24 grams

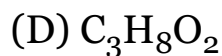
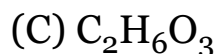
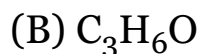
(B) 32 grams

(C) 48 grams

(D) 64 grams

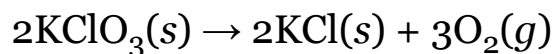
9. A sample of a compound known to consist of only carbon, hydrogen, and oxygen is found to have a total mass of 29.05 g. If the mass of the carbon is 18.02 g and the mass of the hydrogen is 3.03 g, what is the empirical formula of the compound?





Use the following information to answer questions 10-12.

When heated in a closed container in the presence of a catalyst, potassium chlorate decomposes into potassium chloride and oxygen gas via the following reaction:



10. If 12.25 g of potassium chlorate decomposes, how many grams of oxygen gas will be generated?

(A) 1.60 g

(B) 3.20 g

(C) 4.80 g

(D) 18.37 g

11. Approximately how many liters of oxygen gas will be evolved at STP?

(A) 2.24 L

(B) 3.36 L

(C) 4.48 L

(D) 22.4 L

12. If the temperature of the gas is doubled while the volume is held constant, what will happen to the pressure exerted by the gas and why?

- (A) It will also double, because the gas molecules will be moving faster.
- (B) It will also double, because the gas molecules are exerting a greater force on each other.
- (C) It will be cut in half, because the molecules will lose more energy when colliding.
- (D) It will increase by a factor of 4, because the kinetic energy will be four times greater.

13. A sample of a hydrate of CuSO_4 with a mass of 250 grams was heated until all the water was removed. The sample was then weighed and found to have a mass of 160 grams. What is the formula for the hydrate?

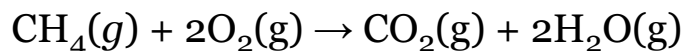
- (A) $\text{CuSO}_4 \cdot 10\text{H}_2\text{O}$
- (B) $\text{CuSO}_4 \cdot 7\text{H}_2\text{O}$
- (C) $\text{CuSO}_4 \cdot 5\text{H}_2\text{O}$
- (D) $\text{CuSO}_4 \cdot 2\text{H}_2\text{O}$

14.
$$\text{CaCO}_3(s) + 2\text{H}^+(aq) \rightarrow \text{Ca}^{2+}(aq) + \text{H}_2\text{O}(l) + \text{CO}_2(g)$$

If the reaction above took place at standard temperature and pressure and 150 grams of $\text{CaCO}_3(s)$ were consumed, what was the volume of $\text{CO}_2(g)$ produced at STP?

- (A) 11 L
- (B) 22 L
- (C) 34 L
- (D) 45 L

15. A gaseous mixture at 25°C contained 1 mole of CH₄ and 2 moles of O₂ and the pressure was measured at 2 atm. The gases then underwent the reaction shown below.



What was the pressure in the container after the reaction had gone to completion and the temperature was allowed to return to 25°C?

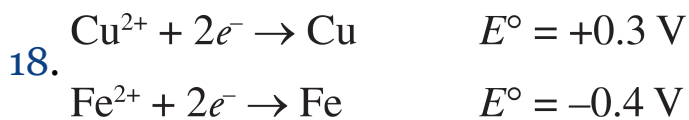
- (A) 1 atm
(B) 2 atm
(C) 3 atm
(D) 4 atm
16. During a chemical reaction, NO(*g*) gets reduced and no nitrogen-containing compound is oxidized. Which of the following is a possible product of this reaction?
- (A) NO₂(*g*)
(B) N₂(*g*)
(C) NO₃⁻(*aq*)
(D) NO₂⁻(*aq*)
17. Which expression below should be used to calculate the mass of copper that can be plated out of a 1.0 *M* Cu(NO₃)₂ solution using a current of 0.75 A for 5.0 minutes?

(A) $\frac{(5.0)(60)(0.75)(63.55)}{(96500)(2)}$

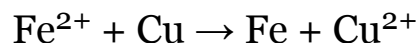
(B) $\frac{(5.0)(60)(63.55)(2)}{(0.75)(96500)}$

$$(C) \frac{(5.0)(60)(96500)(0.75)}{(63.55)(2)}$$

$$(D) \frac{(5.0)(60)(96500)(63.55)}{(0.75)(2)}$$



Based on the reduction potentials given above, what is the reaction potential for the following reaction?



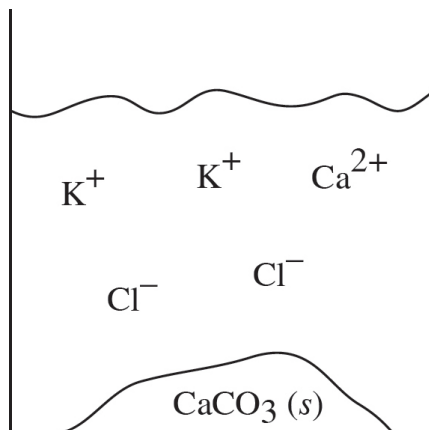
(A) -0.7 V

(B) -0.1 V

(C) $+0.1 \text{ V}$

(D) $+0.7 \text{ V}$

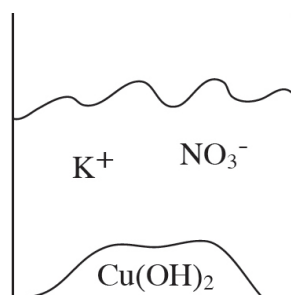
19. Solutions of potassium carbonate and calcium chloride are mixed, and the particulate representation below shows which are present in significant amounts after the reaction has gone to completion.



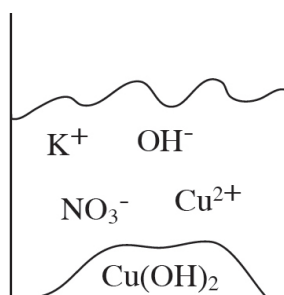
Which of the two original solutions is the limiting reagent and why?

- (A) The potassium carbonate, because of the polyatomic anion
- (B) The potassium carbonate, because there is no carbonate left after the reaction
- (C) The calcium chloride, because there is an excess of calcium ions post-reaction
- (D) The calcium chloride, because the component ions are smaller than those in potassium carbonate

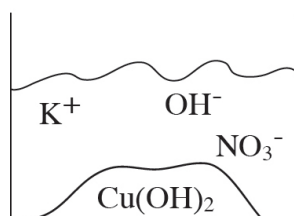
20. A student mixes equimolar amounts of KOH and $\text{Cu}(\text{NO}_3)_2$ in a beaker. Which of the following particulate diagrams correctly shows all species present after the reaction occurs?



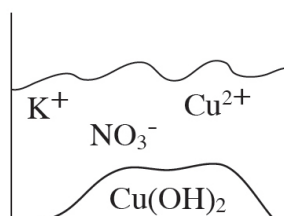
(A)



(B)



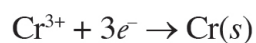
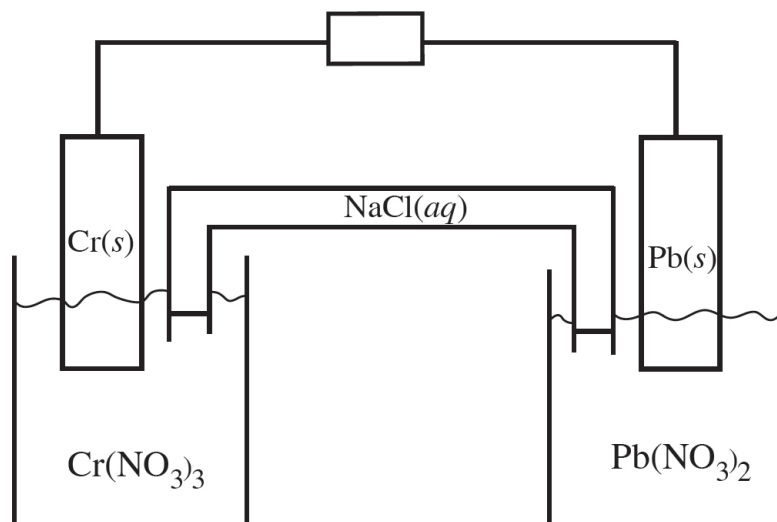
(C)



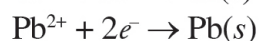
(D)

Use the following information to answer questions 21-23.

A voltaic cell is created using the following half-cells:



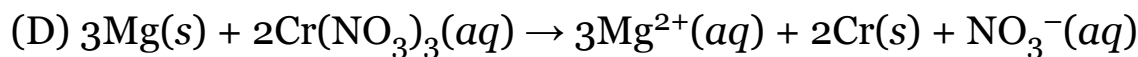
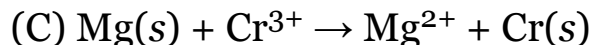
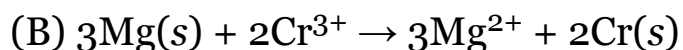
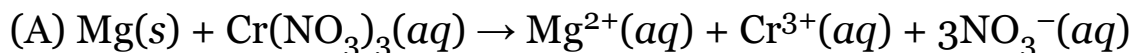
$$E^{\circ} = -0.41 \text{ V}$$



$$E^{\circ} = -0.12 \text{ V}$$

The concentrations of the solutions in each half-cell are 1.0 *M*.

21. Which net ionic equation below represents a possible reaction that takes place when a strip of magnesium metal is oxidized by a solution of chromium (III) nitrate?



22. Which of the following best describes the activity in the salt bridge as the reaction progresses?

(A) Electrons flow through the salt bridge from the Pb/Pb²⁺ half-cell to the Cr/Cr³⁺ half-cell.

(B) Pb²⁺ flows to the Cr/Cr³⁺ half-cell, and Cr³⁺ flows to the Pb/Pb²⁺ half-cell.

(C) Na^+ flows to the Cr/Cr^{3+} half-cell, and Cl^- flows to the Pb/Pb^{2+} half-cell.

(D) Na^+ flows to the Pb/Pb^{2+} half-cell, and Cl^- flows to the Cr/Cr^{3+} half-cell.

23. Based on the given reduction potentials, which of the following would lead to a reaction?

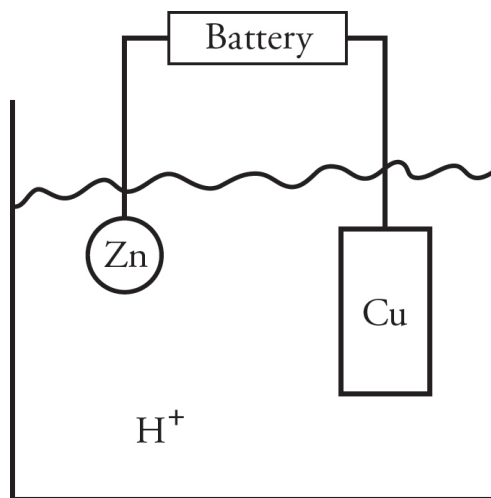
(A) Placing some $\text{Cr}(\text{s})$ in a solution containing Pb^{2+} ions

(B) Placing some $\text{Pb}(\text{s})$ in a solution containing Cr^{3+} ions

(C) Placing some $\text{Cr}(\text{s})$ in a solution containing Cr^{3+} ions

(D) Placing some $\text{Pb}(\text{s})$ in a solution containing Pb^{2+} ions

Use the following information to answer questions 24-26.



Pennies are made primarily of zinc, which is coated with a thin layer of copper through electroplating, using a setup like the one above. The solution in the beaker is a strong acid (which produces H^+ ions), and the cell is wired so that the copper electrode is the anode and zinc penny is the cathode. Use the following reduction potentials to answer questions 24-26.

Half-Reaction	Standard Reduction Potential
---------------	------------------------------

$\text{Cu}^{2+} + 2e \rightarrow \text{Cu}(s)$	+0.34 V
$2\text{H}^+ + 2e \rightarrow \text{H}_2(g)$	0.00 V
$\text{Ni}^{2+} + 2e \rightarrow \text{Ni}(s)$	0.25 V
$\text{Zn}^{2+} + 2e \rightarrow \text{Zn}(s)$	0.76 V

24. When the cell is connected, which of the following reactions takes place at the anode?

- (A) $\text{Cu}^{2+} + 2e^- \rightarrow \text{Cu}(s)$
- (B) $\text{Cu}(s) \rightarrow \text{Cu}^{2+} + 2e$
- (C) $2\text{H}^+ + 2e^- \rightarrow \text{H}_2(g)$
- (D) $\text{H}_2(g) \rightarrow 2\text{H}^+ + 2e$

25. What is the required voltage to make this cell function?

- (A) 0.34 V
- (B) 0.42 V
- (C) 0.76 V
- (D) 1.10 V

26. If, instead of copper, a nickel bar were to be used, could nickel be plated onto the zinc penny effectively? Why or why not?

- (A) Yes, nickel's SRP is greater than that of zinc, which is all that is required for nickel to be reduced at the cathode
- (B) Yes, nickel is able to take electrons from the H^+ ions in solution, allowing it to be reduced
- (C) No, nickel's SRP is lower than that of H^+ ions, which means the only product being produced at the cathode would be

hydrogen gas

(D) No, nickel's SRP is negative, meaning it cannot be reduced in an electrolytic cell

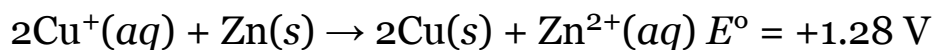
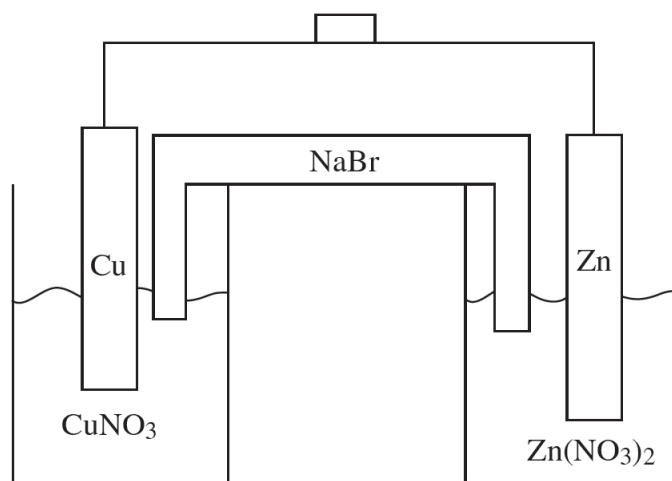
Use the following information to answer questions 27-31.

Two half-cells are set up as follows:

Half-Cell A: Strip of $\text{Cu}(s)$ in $\text{CuNO}_3(aq)$

Half-Cell B: Strip of $\text{Zn}(s)$ in $\text{Zn}(\text{NO}_3)_2(aq)$

When the cells are connected according to the diagram below, the following reaction occurs:



27. Correctly identify the anode and cathode in this reaction as well as where oxidation and reduction are taking place.
- (A) Cu is the anode where oxidation occurs, and Zn is the cathode where reduction occurs.
- (B) Cu is the anode where reduction occurs, and Zn is the cathode where oxidation occurs.

- (C) Zn is the anode where oxidation occurs, and Cu is the cathode where reduction occurs.
- (D) Zn is the anode where reduction occurs, and Cu is the cathode where oxidation occurs.
28. How many moles of electrons must be transferred to create 127 g of copper?
- (A) 1 mole of electrons
- (B) 2 moles of electrons
- (C) 3 moles of electrons
- (D) 4 moles of electrons
29. If the $\text{Cu}^+ + \text{e}^- \rightarrow \text{Cu(s)}$ half-reaction has a standard reduction potential of +0.52 V, what is the standard reduction potential for the $\text{Zn}^{2+} + 2\text{e}^- \rightarrow \text{Zn(s)}$ half-reaction?
- (A) +0.76 V
- (B) -0.76 V
- (C) +0.24 V
- (D) -0.24 V
30. As the reaction progresses, what will happen to the overall voltage of the cell?
- (A) It will increase as $[\text{Zn}^{2+}]$ increases.
- (B) It will increase as $[\text{Cu}^+]$ increases.
- (C) It will decrease as $[\text{Zn}^{2+}]$ increases.
- (D) The voltage will remain constant.
31. What will happen in the salt bridge as the reaction progresses?

- (A) The Na^+ ions will flow to the Cu/Cu^+ half-cell.
- (B) The Br^- ions will flow to the Cu/Cu^+ half-cell.
- (C) Electrons will transfer from the Cu/Cu^+ half-cell to the Zn/Zn^{2+} half-cell.
- (D) Electrons will transfer from the Zn/Zn^{2+} half-cell to the Cu/Cu^+ half-cell.

Free-Response Questions

1. 2.54 g of beryllium chloride are completely dissolved into 50.00 mL of water inside a beaker.

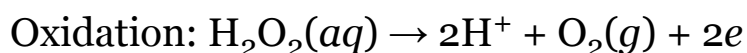
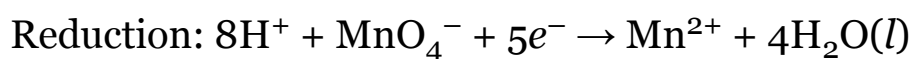
- (a) Draw a particulate representation of all species in the beaker after the solute has dissolved. Your diagram should include at least one beryllium ion, one chloride ion, and four water molecules. Make sure the atoms and ions are correctly sized and oriented relative to each other.
- (b) What is the concentration of beryllium and chloride ions in the beaker?

A solution of 0.850 *M* lead nitrate is then titrated into the beaker, causing a precipitate of lead (II) chloride to form.

- (c) Identify the net ionic reaction occurring in the beaker.
- (d) What volume of lead nitrate must be added to the beaker to cause the maximum precipitate formation?
- (e) What is the theoretical yield of precipitate?
- (f) Students performing this experiment suggested the following techniques to separate the precipitate from the water. Their teacher rejected each idea. Explain why the teacher may have done so, and name the inherent errors of
 - (i) boiling off the water

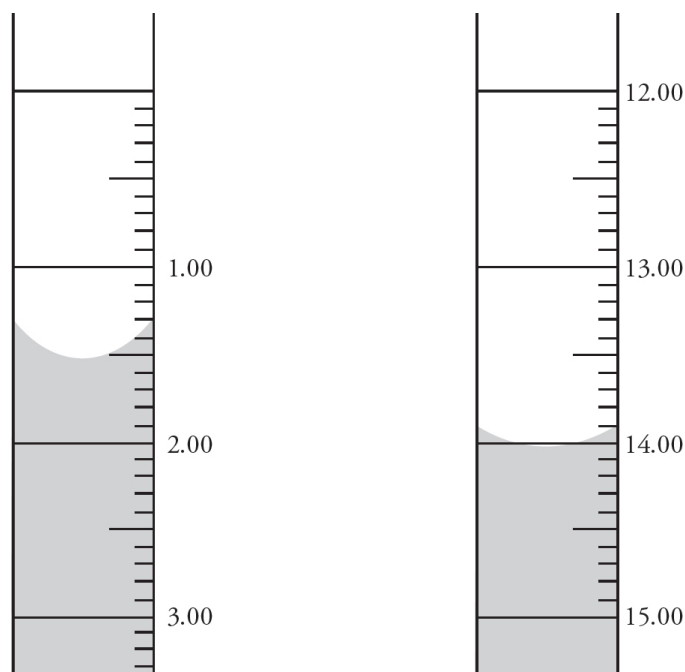
(ii) decanting (pouring off) the water

2. Hydrogen peroxide, H_2O_2 , is a common disinfectant. Pure hydrogen peroxide is a very strong oxidizer, and as such, it is diluted with water to low percentages before being bottled and sold. One method to determine the exact concentration of H_2O_2 in a bottle of hydrogen peroxide is to titrate a sample with a solution of acidified potassium permanganate. This causes the following redox reactions to occur:



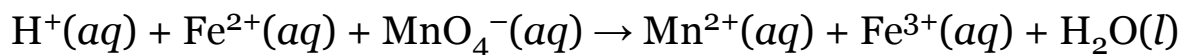
During a titration, a student measures out 5.0 mL of hydrogen peroxide solution into a graduated cylinder, and he pours it into a flask, diluting it to 50.0 mL with water. The student then titrates 0.150 *M* potassium permanganate solution into the flask with constant stirring.

- (a) Write out the full, balanced redox reaction that is taking place during the titration.
- (b) List two observations that the student will see as the titration progresses that are indicative of chemical reactions.



Diagrams of the permanganate in the buret at the start and end of the titration are as follows:

- (c) (i) What volume of KMnO_4 was titrated?
- (ii) What is the concentration of hydrogen peroxide in the original sample?
- (d) How would the precision of the student's results have changed if the hydrogen peroxide sample were measured out in a 50 mL beaker instead of a graduated cylinder?
- (e) How would each of the following errors affect the student's final calculated hydrogen peroxide concentration?
- (i) Not filling the buret tip with solution prior to the titration
- (ii) Not rinsing down the sides of the flask during titration
3. The **unbalanced** reaction between potassium permanganate and acidified iron (II) sulfate is a redox reaction that proceeds as follows:



(a) Provide the equations for both half-reactions that occur below:

(i) Oxidation half-reaction

(ii) Reduction half-reaction

(b) What is the balanced net ionic equation?

A solution of 0.150 *M* potassium permanganate is placed in a buret before being titrated into a flask containing 50.00 mL of iron (II) sulfate solution of unknown concentration. The following data describes the colors of the various ions in solution:

Ion	Color in solution
H ⁺	Colorless
Fe ²⁺	Pale Green
MnO ₄ ⁻	Dark Purple
Mn ²⁺	Colorless
Fe ³⁺	Yellow
K ⁺	Colorless
SO ₄ ²⁻	Colorless

(c) Describe the color of the solution in the flask at the following points:

(i) Before titration begins

(ii) During titration prior to the endpoint

(iii) At the endpoint of the titration

(d) (i) If 15.55 mL of permanganate are added to reach the endpoint, what is the initial concentration of the iron (II) sulfate?

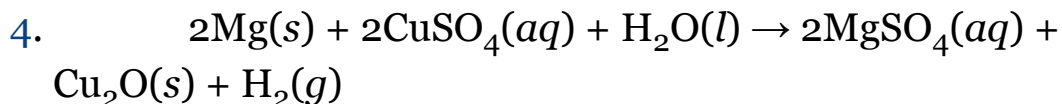
(ii) The actual concentration of the FeSO_4 is 0.250 *M*.

Calculate the percent error.

(e) Could the following errors have led to the experimental result deviating in the direction that it did? You must justify your answers quantitatively.

(i) 55.0 mL of FeSO_4 was added to the flask prior to titration instead of 50.0 mL.

(ii) The concentration of the potassium permanganate was actually 0.160 *M* instead of 0.150 *M*.



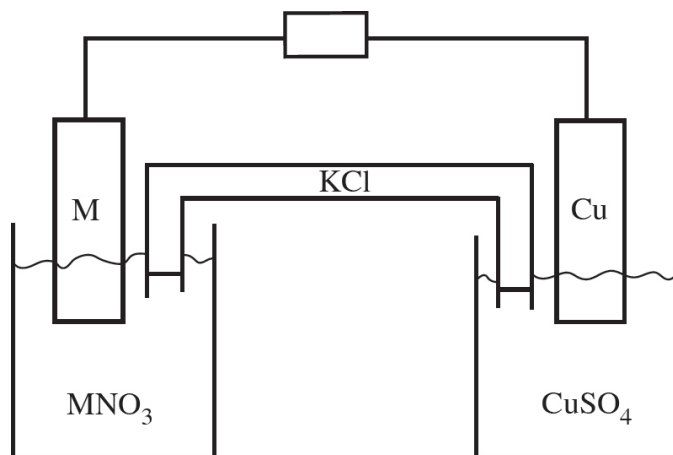
(a) If 1.46 grams of $\text{Mg}(s)$ are added to 500 milliliters of a 0.200-molar solution of CuSO_4 , what is the maximum molar yield of $\text{H}_2(g)$?

(b) When all of the limiting reagent has been consumed in (a), how many moles of the other reactant (not water) remain?

(c) What is the mass of the Cu_2O produced in (a)?

(d) What is the value of $[\text{Mg}^{2+}]$ in the solution at the end of the experiment? (Assume that the volume of the solution remains unchanged.)

5. A student performs an experiment in which a bar of unknown metal M is placed in a solution with the formula MNO_3 . The metal is then hooked up to a copper bar in a solution of CuSO_4 as shown below. A salt bridge that contains aqueous KCl links the cell together.



The cell potential is found to be +0.74 V. Separately, when a bar of metal M is placed in the copper sulfate solution, solid copper starts to form on the bar. When a bar of copper is placed in the MNO_3 solution, no visible reaction occurs.

The following gives some reduction potentials for copper:

Half-reaction	E
$\text{Cu}^{2+} + 2e^- \rightarrow \text{Cu}(s)$	0.34 V
$\text{Cu}^{2+} + e^- \rightarrow \text{Cu}^+$	0.15 V
$\text{Cu}^+ + e^- \rightarrow \text{Cu}(s)$	0.52 V

- Write the net ionic equation that takes place in the Cu/M cell.
- What is the standard reduction potential for metal M?
- Which metal acted as the anode and which as the cathode?
Justify your answer.
- On the diagram of the cell, indicate which way the electrons are flowing in the wire. Additionally, indicate any ionic movement occurring in the salt bridge.
- What would happen to the voltage of the reaction in the Cu/M cell if the concentration of the CuSO_4 increased while

the concentration of the MNO_3 remained constant? Justify your answer.

6. Two electrodes are inserted into a solution of nickel (II) fluoride and a current of 2.20 A is run through them. A list of standard reduction potentials is as follows:

Half-reaction	E
$\text{O}_2(g) + 4\text{H}^+ + 4e^- \rightarrow \text{H}_2\text{O}(l)$	1.23 V
$\text{F}_2(g) + 2e^- \rightarrow 2\text{F}^-$	2.87 V
$2\text{H}_2\text{O}(l) + 2e^- \rightarrow \text{H}_2(g) + 2\text{OH}^-$	-0.83 V
$\text{Ni}^{2+} + 2e^- \rightarrow \text{Ni}(s)$	-0.25 V

- (a) Write the net ionic equation that takes place during this reaction.
- (b) Qualitatively describe what an observer would see taking place at each electrode.
- (c) Will the solution become acidic, basic, or remain neutral as the reaction progresses?
- (d) How long would it take to create 1.2 g of $\text{Ni}(s)$ at the cathode?

CHAPTER 5 ANSWERS AND EXPLANATIONS

Multiple-Choice

1. **B**

Carbonates are insoluble when paired with iron. Iron has a charge of +3 and carbonate has a charge of -2 . To cancel out, both charges need to have a magnitude of 6, requiring two iron atoms and three carbonate ions. The best representation of that is (B).

2. **D**

When those solutions mix, a precipitate of AgCl will form, removing those ions from the solution. The remaining ions (K^+ and NO_3^-) do not react and remain the same as they were when they started.

3. **D**

The precipitate that would form is PbCl_2 , meaning that both NO_3^- and Na^+ are spectator ions. The precipitate would require twice as many chloride ions as lead ions. Therefore, if equal moles of both are used, the NaCl would be the limiting reagent, and almost all of the chloride ions would be present in the solid, with very few left in the solution. There would, however, still be significant Pb^{2+} remaining in solution, as it is in excess.

4. **C**

Via the information given from the solubility rules, you can determine that the precipitate would be copper (I) carbonate, which has a formula of Cu_2CO_3 . In the net ionic

equation, spectator ions (in this case, K^+ and Cl^-) cancel out and do not appear in the final reaction.

5. **C**

Z^{2+} was able to reduce to solid Z by taking electrons from metal X, so Z must have a higher reduction potential than X. Y^{2+} was unable to take electrons from metal X, and therefore Y must have a lower reduction potential than X.

6. **A**

In (D), chromium is a pure element and has an oxidation number of 0. In (C), chromium's oxidation number is equal to its charge of +3, and in (B), it must balance the -2 on the oxygen, so it has a charge of +2. In (A), the total charge on the ion is -2 , and each oxygen is -2 . Solving the following, where X is the oxidation number on chromium: $X + -2(4) = -2$. So, the oxidation number is +6.

7. **C**

For any reaction (exothermic or endothermic), the energy level of the activated complex is always above the energy level of both the products and reactants.

8. **D**

$$\text{Moles} = \frac{\text{grams}}{\text{MW}}$$

$$\text{Moles of calcium hydroxide} = \frac{(148 \text{ g})}{(74 \text{ g/mol})} = 2 \text{ moles}$$

Every mole of $Ca(OH)_2$ contains 2 moles of oxygen.

So there are $(2)(2) = 4$ moles of oxygen

Grams = (moles)(MW)

So grams of oxygen = (4 mol)(16 g/mol) = 64 grams

9. **B**

First, the mass of the oxygen must be calculated: $29.05 \text{ g} - 18.02 \text{ g} - 3.03 \text{ g} = 8.00 \text{ g}$.

Converting each of those to moles yields 0.5 moles of oxygen, 1.5 moles of carbon, and 3.0 moles of hydrogen.

Thus, for every one oxygen atom there are three carbon atoms and six hydrogen atoms.

10. **C**

$$12.25 \text{ g KClO}_3 \times \frac{1 \text{ mol KClO}_3}{122.25 \text{ g KClO}_3} = 0.1000 \text{ mol KClO}_3$$

$$0.1000 \text{ mol KClO}_3 \times \frac{3 \text{ mol O}_2}{2 \text{ mol KClO}_3} = 0.1500 \text{ mol O}_2$$

$$0.1500 \text{ mol O}_2 \times \frac{32.00 \text{ g O}_2}{1 \text{ mol O}_2} = 4.80 \text{ g O}_2$$

Note that even without a calculator, you are expected to be able to do simple calculations such as the ones above.

11. **B**

$$0.1500 \text{ mol O}_2 \times \frac{22.4 \text{ L}}{1 \text{ mol O}_2} = 3.36 \text{ L O}_2$$

12. **A**

Pressure is a measure of the amount of force with which the gas particles are hitting the container walls. An increase in temperature is indicative of increased molecular velocity. If the molecules are moving faster, they will not only have more energy when they hit the container walls, but they will also hit those walls more often. Both of those factors contribute to the increased pressure.

13. C

The molecular weight of CuSO_4 is 160 g/mol, so you have only 1 mole of the hydrate. The lost mass was due to water, so 1 mole of the hydrate must have contained 90 grams of H_2O .

$$\text{Moles} = \frac{\text{grams}}{\text{MW}}$$

$$\text{Moles of water} = \frac{(90 \text{ g})}{(18 \text{ g/mol})} = 5 \text{ moles}$$

So if 1 mole of hydrate contains 5 moles of H_2O , then the formula for the hydrate must be $\text{CuSO}_4 \cdot 5 \text{ H}_2\text{O}$.

14. C

$$\text{Moles} = \frac{\text{grams}}{\text{MW}}$$

$$\text{Moles of CaCO}_3 = \frac{(150 \text{ g})}{(100 \text{ g/mol})} = 1.5 \text{ moles}$$

From the balanced equation, for every mole of CaCO_3 consumed, one mole of CO_2 is produced. So 1.5 moles of CO_2 are produced.

At STP, volume of gas = (moles)(22.4 L)

So the volume of $\text{CO}_2 = (1.5)(22.4) = 34 \text{ L}$

15. **B**

All of the reactants are consumed in the reaction and the temperature doesn't change, so the pressure will change only if the number of moles of gas changes over the course of the reaction. The number of moles of gas (3 moles) doesn't change in the balanced equation, so the pressure will remain the same (2 atm) at the end of the reaction as at the beginning.

16. **B**

The oxidation state of nitrogen in NO is +2. If the nitrogen is reduced, that value must get more negative. The oxidation state of nitrogen in N_2 is 0, so that fits the bill. The oxidation state on the nitrogen in the other choices is greater than +2.

17. **A**

$$5.0 \text{ min} \times \frac{60.0 \text{ s}}{1.0 \text{ min}} \times \frac{0.75 \text{ C}}{1.0 \text{ s}} \times \frac{1 \text{ mol } e^-}{96,500 \text{ C}} \\ \times \frac{1 \text{ mol Cu}}{2 \text{ mol } e^-} \times \frac{63.55 \text{ g Cu}}{1 \text{ mol Cu}}$$

18. **A**

Add the reduction potential for Fe^{2+} (-0.4 V) to the oxidation potential for Cu (-0.3 V , the reverse of the reduction potential) to get -0.7 V .

19. **B**

If extra Ca^{2+} ions are in solution, that means there were not enough CO_3^{2-} ions present for the Ca^{2+} ions to fully react.

20. **D**

The overall reaction (excluding spectator ions) is: $2\text{OH}^- + \text{Cu}^{2+} \rightarrow \text{Cu}(\text{OH})_2(\text{s})$. Both the K^+ and the NO_3^- are spectator ions which are present in the solution both before and after the reaction. Additionally, if equimolar amounts of the two reactants are initially present, the OH^- will run out before the Cu^{2+} , meaning that some Cu^{2+} ions will also be present in the final solution.

21. **B**

Cr^{3+} needs to gain three electrons to reduce into $\text{Cr}(\text{s})$, while $\text{Mg}(\text{s})$ loses 2 electrons to oxidize into Mg^{2+} . To balance the electrons, the reduction half-reaction must be multiplied by two and the oxidation half-reaction must be multiplied by three in order to have six electrons on each side. The nitrate ion is a spectator and would not appear in the net ionic equation.

22. **D**

As the reaction progresses, Pb^{2+} is being reduced to create $\text{Pb}(\text{s})$. This decreases the amount of positive charge in the solution, which is replaced by the Na^+ from the salt bridge. At the anode, excess Cr^{3+} is being created, which is balanced by the Cl^- anions from the salt bridge.

23. **A**

For a reaction to occur, the reduction potential of the ions in solution must exceed the reduction potential for the ions of the corresponding metal. If $\text{Cr}(\text{s})$ were placed in a

solution containing Pb^{2+} , the Pb^{2+} would take electrons from the Cr(s) and create Pb(s) and Cr^{3+} .

24. **B**

Oxidation occurs at the anode, and that entails the loss of electrons. As there is no hydrogen gas present in the solution to start, the only substance that can be reduced at the anode is the solid copper itself.

25. **A**

The half-reaction occurring at the anode is $2\text{H}^+ + 2e^- \rightarrow \text{H}_2(g)$, as the only substance present at the beginning of the reaction can that can be reduced is the hydrogen ion (initially, there are no Zn^{2+} or Cu^{2+} ions in solution). With copper is being oxidized at the anode, the calculation then becomes $(0.00 \text{ V}) + (-0.34 \text{ V}) = -0.34 \text{ V}$. That means a voltage differential of 0.34 V will be required for this reaction to occur.

26. **C**

In the initial Cu/Zn cell, as the electrolysis proceeds Cu^{2+} ions are produced and become part of the solution. The SRP of those ions is greater than that the SRP of H^+ ions, so later in the reaction, the Cu^{2+} is reduced into solid copper at the zinc cathode—this is what causes the copper to plate out on the zinc. If nickel were used as the anode, Ni^{2+} would be in solution, however, H^+ has a higher reduction potential than Ni^{2+} , which means the H^+ reduction would continue to occur at the cathode, and no nickel would be plated out of solution.

27. **C**

The oxidation state of copper changes from +1 to 0, meaning it has gained electrons and is being reduced, and reduction occurs at the cathode. Zinc's oxidation state changes from 0 to +2, meaning it has lost electrons and is being oxidized, which occurs at the anode.

28. **B**

127 g is equal to 2 moles of copper, which is what appears on the balanced equation. To change one mole of copper from +1 to 0, 1 mole of electrons is required. Twice as many moles being created means twice as many electrons are needed.

29. **B**

$$E_{\text{cell}} = E_{\text{red}} + E_{\text{ox}}$$

$$1.28 \text{ V} = 0.52 \text{ V} + E_{\text{ox}}$$

$$E_{\text{ox}} = 0.76 \text{ V}$$

$$-E_{\text{ox}} = E_{\text{red}}$$

$$E_{\text{red}} = -0.76 \text{ V}$$

30. **C**

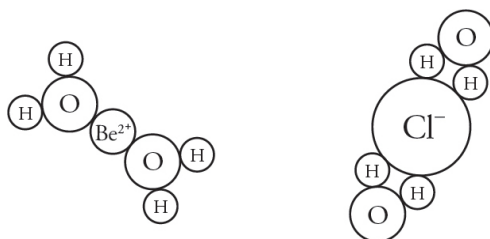
As the reaction progresses, $[\text{Cu}^+]$ will decrease and $[\text{Zn}^{2+}]$ will increase. With a lower concentration on the reactants side and a higher concentration on the products side, the reaction will shift left, decreasing the overall potential of the reaction. Doing this will increase the value of Q ($Q = [\text{Zn}^{2+}]/[\text{Cu}^+]$), bringing the reaction closer to equilibrium and decreasing the overall potential of the reaction.

31. A

The electron transfer does not happen across the salt bridge, eliminating (C) and (D). As the reaction progresses and $[\text{Cu}^+]$ decreases in the copper half-cell, positively charged sodium ions are transferred in to keep the charge balanced within the half-cell.

Free-Response

1. (a) The beryllium ions only have one full energy level, while chloride ions have three and would thus be much larger. The positive beryllium ions would be attracted to the negative dipoles of the water molecules, and the negative chloride ions would be attracted to the positive dipoles of the water molecules.

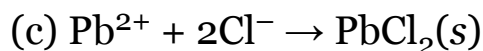


- (b) First, calculate the concentration of the BeCl_2 .

$$2.54 \text{ g BeCl}_2 \times \frac{1 \text{ mol BeCl}_2}{79.91 \text{ g BeCl}_2} \times \frac{0.0317 \text{ mol}}{0.0500 \text{ L}} = 0.635 M$$

BeCl_2

For every mole of BeCl_2 that dissociates, one Be^{2+} and two Cl^- ions are produced. Thus, the concentration of the Be^{2+} will be the same as the base concentration of the BeCl_2 , and the concentration of the Cl^- will be twice that. So, $[\text{Be}^{2+}] = 0.635 M$ and $[\text{Cl}^-] = 1.27 M$.



- (d) There must be one lead ion present for every two chloride ions in order to create the maximum amount of precipitate. Figure out the number of moles of chloride from the concentration and volume, which you already have, and then convert:

$$1.27 \text{ M Cl}^- = \frac{n}{0.0500 \text{ L}}$$

$$n = 0.0635 \text{ mol Cl}^- \times \frac{1 \text{ mol Pb}^{2+}}{2 \text{ mol Cl}^-} = 0.0318 \text{ mol Pb}^{2+}$$

From there, you can use the given concentration of the lead nitrate solution to determine how much of it is required.

$$0.850 \text{ M} = \frac{0.0318 \text{ mol}}{V}$$

$$V = 0.0374 \text{ L} = 37.4 \text{ mL}$$

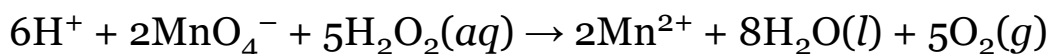
- (e) If both reactants are mixed in equal amounts, either value can be used to calculate the amount of precipitate that will be created.

$$0.0318 \text{ mol Pb}^{2+} \times \frac{1 \text{ mol PbCl}_2}{1 \text{ mol Pb}^{2+}} \times \frac{278.1 \text{ g}}{1 \text{ mol PbCl}_2} = 8.84 \text{ g}$$

- (f) (i) Boiling the water would evaporate all the water, but it would also cause the spectator ions to no longer be dissolved in water. In addition to any PbCl_2 that was created, the excess beryllium and nitrate ions would adhere to the solid, giving a falsely high molar mass and an impure product.

(ii) While decanting the solution, smaller particles of precipitate may get poured out with the water. Decanting the solution into a filtered funnel setup would be a much better option, as the filter paper would then catch all of the precipitate while allowing the water (which would carry the spectator ions) through.

2. (a) In a full redox reaction, the electrons from each half-reaction must cancel. To do that, the reduction reaction must be multiplied by two, and the oxidation reaction multiplied by five. When combining the reactions after doing so, this yields:



Note that 10 of the hydrogen ions cancel out when the reactions combine.

- (b) A gas will be evolved as the hydrogen peroxide oxidizes into oxygen gas. Additionally, there will be a color change as the purple permanganate solution reduces into colorless manganese ions.

- (c) (i) $14.03 \text{ mL} - 1.52 \text{ mL} = 12.51 \text{ mL}$

- (ii) First, calculate the moles of permanganate that were titrated.

$$0.150 \text{ M} = \frac{n}{0.01251 \text{ L}}$$

$$n = 0.00188 \text{ mol MnO}_4^-$$

Then use the stoichiometric ratios from the balanced equation to convert to moles of hydrogen peroxide.

$$0.00188 \text{ mol MnO}_4^- \times \frac{5 \text{ mol H}_2\text{O}_2}{2 \text{ mol MnO}_4^-} = 0.00470 \text{ mol H}_2\text{O}_2$$

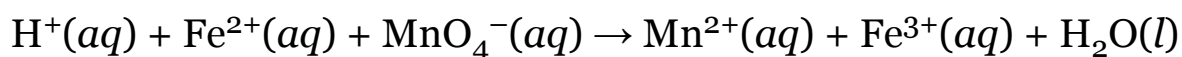
Finally, convert that to the concentration of H_2O_2 . We will use a volume of 5.00 mL and not 50.0 mL in our calculations because we are interested in the concentration of the solution before it was diluted with water.

$$\frac{0.00470 \text{ mol H}_2\text{O}_2}{0.0050 \text{ L H}_2\text{O}_2} = 0.94 \text{ M H}_2\text{O}_2 = 0.94 \text{ M H}_2\text{O}_2$$

- (d) Beakers are precise to the ones places; you cannot accurately measure out a volume in a beaker to any number of decimal places. Graduated cylinders are accurate to one decimal place, so using a beaker would reduce the number of significant figures in your answer by one.
- (e) (i) If the tip of the buret is not filled prior to dispensing the permanganate solution from it, the first milliliter or so that would be dispensed is air instead of solution. Thus, the experimental volume of permanganate would be artificially high (instead of 12.51 mL of permanganate, it would have really been about 11.5 mL of permanganate with the rest being air). If the recorded volume of the permanganate is too high, that will eventually lead to a calculated $[\text{H}_2\text{O}_2]$ that is also too high.
- (ii) During a titration, it is best practice to regularly rinse down the sides of the flask you are titrating into, just in case any of the titrated solution splashes onto it and does not mix with the solution you are titrating into. If you do not do this, some of the permanganate that leaves the

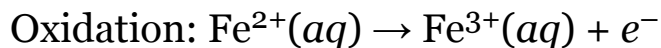
buret might stick to the sides of the flask and not react with the hydrogen peroxide. This would mean the recorded volume of permanganate is too high (not all of the measured permanganate actually reacted), which would again lead to an artificially high calculate concentration of hydrogen peroxide.

3. The unbalanced reaction between potassium permanganate and acidified iron (II) sulfate is a redox reaction that proceeds as follows:

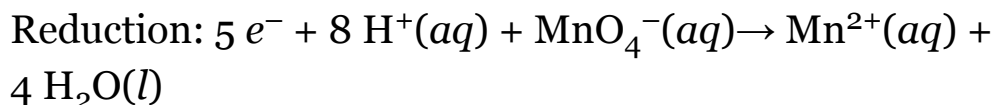


- (a) Provide the equations for both half-reactions that occur below:

- (i) Oxidation half-reaction:

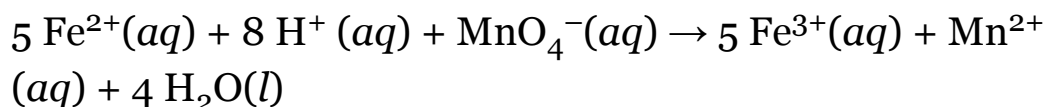


- (ii) Reduction half-reaction:



- (b) What is the balanced net ionic equation?

The oxidation reaction must be multiplied by a factor of 5 in order for the electrons to balance out. So:



A solution of 0.150 M potassium permanganate, a dark purple solution, is placed in a buret before being titrated into a flask containing 50.00 mL of iron (II) sulfate

solution of unknown concentration. The following data describes the colors of the various ions in solution:

Ion	Color in solution
H^+	Colorless
Fe^{2+}	Pale Green
MnO_4^-	Dark Purple
Mn^{2+}	Colorless
Fe^{3+}	Yellow
K^+	Colorless
SO_4^{2-}	Colorless

(c) Describe the color of the solution in the flask at the following points:

(i) Before titration begins

The only ions present in the flask are Fe^{2+} , SO_4^{2-} , and H^+ . The latter two are colorless, so the solution would be pale green.

(ii) During titration prior to the endpoint

The MnO_4^- is reduced to Mn^{2+} upon entering the flask, and the Fe^{2+} ions are oxidized into Fe^{3+} ions. The solution would become less green as more yellow as more Fe^{3+} ions are formed, as all other ions present are colorless.

(iii) At the endpoint of the titration

After the Fe^{2+} ions have all been oxidized, there is nothing left to donate electrons to the MnO_4^- ions. Therefore, they will no longer be reduced upon entering the flask, and the solution will take on a light purplish/yellow hue due to the mixture of MnO_4^- and Fe^{3+} ions.

- (d) (i) If 15.55 mL of permanganate are added to reach the endpoint, what is the initial concentration of the iron (II) sulfate?

First, the moles of permanganate added must be calculated:

$$0.150\text{ M} = \frac{n}{0.0155\text{ L}}$$

$$n = 2.33 \times 10^{-3}\text{ mol MnO}_4^-$$

Then, the moles of iron (II) can be determined via stoichiometry:

$$2.33 \times 10^{-3}\text{ mol MnO}_4^- \times \frac{5\text{ mol Fe}^{2+}}{1\text{ mol MnO}_4^-} = 0.0117\text{ mol Fe}^{2+}$$

Finally, the concentration of the FeSO_4 can be determined:

$$\text{Molarity} = \frac{0.0117\text{ mol}}{0.050\text{ L}} = 0.234\text{ M}$$

- (ii) The actual concentration of the FeSO_4 is 0.250 M. Calculate the percent error.

$$\text{Percent error} = \frac{\text{actual value} - \text{experimental value}}{\text{actual value}} \times 100$$

$$\text{Percent error} = \frac{0.250 - 0.234}{0.250} \times 100 = 6.40\% \text{ error}$$

(e) Could the following errors have led to the experimental result deviating in the direction that it did? You must justify your answers quantitatively.

- (i) 55.0 mL of FeSO_4 was added to the flask prior to titration instead of 50.0 mL.

If the volume of FeSO_4 was artificially low in the calculations, that would lead to the experimental value for the concentration of FeSO_4 being artificially high. As the calculated value for the concentration of FeSO_4 was too low, this error source is not supported by data.

- (ii) The concentration of the potassium permanganate was actually 0.160 *M* instead of 0.150 *M*.

If the molarity of the permanganate was artificially low in the calculations, the moles of permanganate, and by extension, the moles of Fe^{2+} would also be artificially low. This would lead to an artificially low value for the concentration of FeSO_4 . This matches with the experimental results and is thus supported by the data.

4. (a) You need to find the limiting reagent. There's plenty of water, so it must be one of the other two reactants.

$$\text{Moles} = \frac{\text{grams}}{\text{MW}}$$

$$\text{Moles of Mg} = \frac{(1.46 \text{ g})}{(24.3 \text{ g/mol})} = 0.060 \text{ moles}$$

$$\text{Moles} = (\text{molarity})(\text{volume})$$

$$\text{Moles of CuSO}_4 = (0.200 \text{ M})(0.500 \text{ L}) = 0.100 \text{ moles}$$

From the balanced equation, Mg and CuSO₄ are consumed in a 1:1 ratio, so run out of Mg first. Mg is the limiting reagent, and then use it to find the yield of H₂.

From the balanced equation, 1 mole of H₂ is produced for every 2 moles of Mg consumed, so the number of moles of H₂ produced will be half the number of moles of Mg consumed.

$$\text{Moles of H}_2 = \frac{1}{2}(0.060 \text{ mol}) = 0.030 \text{ moles}$$

(b) Mg is the limiting reagent, so some CuSO₄ will remain.

From the balanced equation, Mg and CuSO₄ are consumed in a 1:1 ratio, so when 0.060 moles of Mg are consumed, 0.060 moles of CuSO₄ are also consumed.

$$\text{Moles of CuSO}_4 \text{ remaining} = (0.100 \text{ mol}) - (0.060 \text{ mol}) = 0.040 \text{ moles}$$

(c) From the balanced equation, 1 mole of Cu₂O is produced for every 2 moles of Mg consumed, so the number of moles of Cu₂O produced will be half the number of moles of Mg consumed.

$$\text{Moles of Cu}_2\text{O} = \frac{1}{2}(0.060 \text{ mol}) = 0.030 \text{ moles}$$

$$\text{Grams} = (\text{moles})(\text{MW})$$

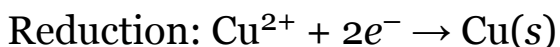
$$\text{Grams of Cu}_2\text{O} = (0.030 \text{ mol})(143 \text{ g/mol}) = 4.29 \text{ grams}$$

(d) All of the Mg consumed ends up as Mg^{2+} ions in the solution.

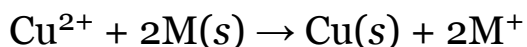
$$\text{Molarity} = \frac{\text{moles}}{\text{liters}}$$

$$[\text{Mg}^{2+}] = \frac{(0.060 \text{ mol})}{(0.500 \text{ L})} = 0.120 \text{ M}$$

5. (a) When metal M was placed in the copper solution, a reaction occurred. Therefore, the copper must have the higher reduction potential and thus is reduced in the Cu/M cell. The half-reactions are:



The oxidation half-reaction must be multiplied by two to balance the electrons before combining the reactions to yield:



$$(b) E_{\text{red}} + E_{\text{ox}} = +0.74 \text{ V}$$

The reduction potential for $\text{Cu}^{2+} + 2e^- \rightarrow \text{Cu(s)}$ is known:

$$0.34 \text{ V} + E_{\text{ox}} = +0.74 \text{ V}$$

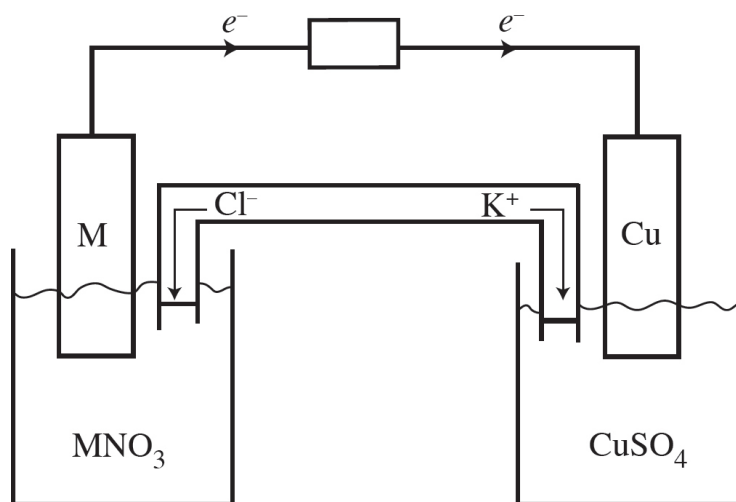
$$E_{\text{ox}} = +0.40 \text{ V}$$

The reduction potential for metal M is the opposite of its oxidation potential.

$$E_{\text{red}} = -0.40 \text{ V}$$

(c) Reduction occurs at the cathode, so copper is the cathode. Oxidation occurs at the anode, so M is the anode.

(d)

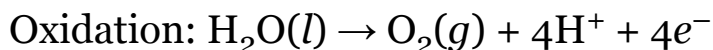
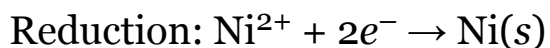


Electrons should be flowing from metal M to the copper bar. In the salt bridge, the K^+ ions will flow toward the copper solution to replace the Cu^{2+} being reduced into Cu(s) . The Cl^- will flow toward the solution for metal M in order to balance out the charge of the extra M^+ ions being created via the oxidation of M(s) .

(e) If the concentration of the Cu^{2+} increases, it will cause a shift to the right in accordance with Le Châtelier's principle. This will increase the cell's overall voltage.

6. (a) The two potential reduction reactions are the bottom two, as their reactants (H_2O and Ni^{2+}) are actually present at the start of the reaction. Of the two, the nickel reduction is more positive, and thus will take place. The top two reactions must be flipped to have the reactants (H_2O and F^-) on the reactants side, which also flips the sign. As a

result, the water oxidation has the more positive value (–1.23 as opposed to –2.87) and will therefore occur.



After multiplying the reduction half-reaction by 2 to balance the electrons and combining both half-reactions, the net cell reaction is:



- (b) At the cathode, solid nickel would begin to plate out of solution. At the anode, oxygen gas would form and bubble up to the surface.
- (c) The solution will become more acidic due to the creation of hydrogen ions at the anode.
- (d) $1.20 \text{ g Ni} \times \frac{1 \text{ mol Ni}}{58.69 \text{ g Ni}} \times \frac{2 \text{ mol } e^{-}}{1 \text{ mol N}} \times \frac{96,500 \text{ C}}{1 \text{ mol } e^{-}} \times \frac{1.0 \text{ sec}}{2.20 \text{ C}} =$
1790 seconds

Chapter 6

Big Idea #4: Chemical Reactions and Their Rates

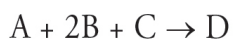
Rates of chemical reactions are determined by details of the molecular collisions.

RATE LAW USING INITIAL CONCENTRATIONS

The rate law for a reaction describes the dependence of the initial rate of a reaction on the concentrations of its reactants. It includes the Arrhenius constant, k , which takes into account the activation energy for the reaction and the temperature at which the reaction occurs. The rate of a reaction is described in terms of the rate of appearance of a product or the rate of disappearance of a reactant. The rate law for a reaction cannot be determined from a balanced equation; it must be determined from experimental data, which is presented on the test in table form.

Here's How It's Done

The data below were collected for the following hypothetical reaction:



Experiment	Initial Concentration of Reactants (M)			Initial Rate of Formation of D (M/sec)
	[A]	[B]	[C]	
1	0.10	0.10	0.10	0.01
2	0.10	0.10	0.20	0.01
3	0.10	0.20	0.10	0.02
4	0.20	0.20	0.10	0.08

The rate law always takes the following form, using the concentrations of the reactants:

$$\text{Rate} = k[A]^x[B]^y[C]^z$$

The greater the value of a reactant's exponent, the more a change in the concentration of that reactant will affect the rate of the reaction. To find the values for the exponents x , y , and z , we need to examine how changes in the individual reactants affect the rate. The easiest way to find the exponents is to see what happens to the rate when the concentration of an individual reactant is doubled.

Let's Look at [A]

From experiment 3 to experiment 4, [A] doubles while the other reactant concentrations remain constant. For this reason, it is useful to use the rate values from these two experiments to calculate x (the order of the reaction with respect to reactant A).

As you can see from the table, the rate quadruples from experiment 3 to experiment 4, going from 0.02 M/sec to 0.08 M/sec.

We need to find a value for the exponent x that relates the doubling of the concentration to the quadrupling of the rate. The value of x can be calculated as follows:

$$(2)^x = 4, \text{ so } x = 2$$

Because the value of x is 2, the reaction is said to be second order with respect to A.

$$\text{Rate} = k[\text{A}]^2[\text{B}]^y[\text{C}]^z$$

Let's Look at [B]

From experiment 1 to experiment 3, [B] doubles while the other reactant concentrations remain constant. For this reason it is useful to use the rate values from these two experiments to calculate y (the order of the reaction with respect to reactant B).

As you can see from the table, the rate doubles from experiment 1 to experiment 3, going from 0.01 M/sec to 0.02 M/sec.

We need to find a value for the exponent y that relates the doubling of the concentration to the doubling of the rate. The value of y can be calculated as follows:

$$(2)^y = 2, \text{ so } y = 1$$

Because the value of y is 1, the reaction is said to be first order with respect to B.

$$\text{Rate} = k[\text{A}]^2[\text{B}][\text{C}]^z$$

Let's Look at [C]

From experiment 1 to experiment 2, [C] doubles while the other reactant concentrations remain constant.

The rate remains the same at 0.01 *M*.

The rate change is $(2)^z = 1$, so $z = 0$.

Because the value of z is 0, the reaction is said to be zero order with respect to C.

$$\text{Rate} = k[\text{A}]^2[\text{B}]$$

Because the sum of the exponents is 3, the reaction is said to be third order overall.

Once the rate law has been determined, the value of the rate constant can be calculated using any of the lines of data on the table. The units of the rate constant are dependent on the order of the reaction, so it's important to carry along units throughout all rate constant calculations.

Let's use experiment 3.

$$k = \frac{\text{Rate}}{[\text{A}]^2[\text{B}]} = \frac{(0.02 \text{ M} / \text{sec})}{(0.10 \text{ M})^2 (0.20 \text{ M})} = 10 \left(\frac{(\text{M})}{(\text{M})^3 (\text{sec})} \right) = 10 \text{ M}^{-2} \cdot \text{sec}^{-1}$$

You should note that we can tell from the coefficients in the original balanced equation that the rate of appearance of D is equal to the rate of disappearance of A and C because the coefficients of all three are the same. The coefficient of D is half as large as the coefficient of B, however, so the rate at which D appears is half the rate at which B disappears.

RATE LAW USING CONCENTRATION AND TIME

It's also useful to have rate laws that relate the rate constant k to the way that concentrations change over time. The rate laws will be different depending on whether the reaction is first, second, or zero order, but each rate law can be expressed as a graph that relates the rate constant, the concentration of a reactant, and the elapsed time.

Zero-Order Rate Laws

The rate of a zero-order reaction does not depend on the concentration of reactants at all, so the rate of a zero-order reaction will always be the same at a given temperature.

$$\text{Rate} = k$$

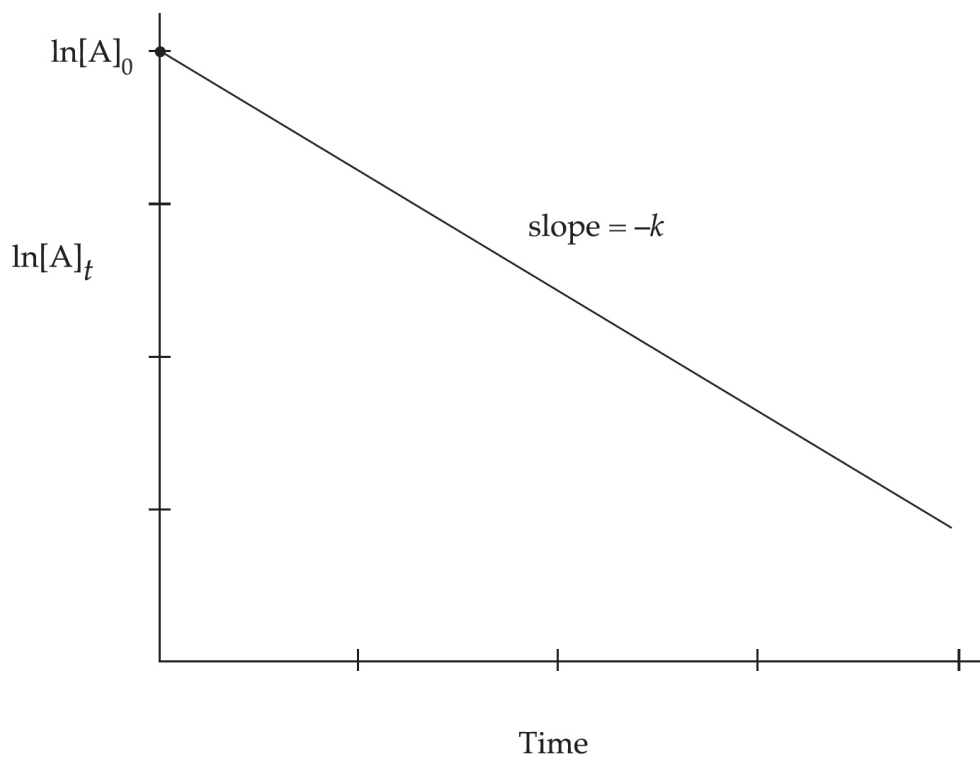
The graph of the change in concentration of a reactant of a zero-order reaction versus time will be a straight line with a slope equal to $-k$.

First-Order Rate Laws

The rate of a first-order reaction depends on the concentration of a single reactant raised to the first power.

$$\text{Rate} = k[A]$$

The rate law for a first-order reaction uses natural logarithms. This means that the use of natural logarithms in the rate law creates a linear graph comparing concentration and time. The use of natural logarithms in the rate law creates a linear graph comparing concentration and time. The slope of the line is given by $-k$ and the y -intercept is given by $\ln[A]_0$.



Using slope-intercept form, we can interpret this graph to come up with a useful equation.

$$y = mx + b$$

$$y = \ln[A]_t$$

$$m = -k$$

$$x = \text{time}$$

$$b = \ln [A]_0$$

After substitution and rearrangement, the slope-intercept equation becomes:

First-Order Rate Law

$$\ln[A]_t = -kt + \ln[A]_0$$

$[A]_t$ = concentration of reactant A at time t

$[A]_0$ = initial concentration of reactant A

k = rate constant

t = time elapsed

Second-Order Rate Laws

The rate of a second-order reaction depends on the concentration of a single reactant raised to the second power.

$$\text{Rate} = k[A]^2$$

The rate law for a second-order reaction uses the inverses of the concentrations.

Second-Order Rate Law

$$\frac{1}{[A]_t} = kt + \frac{1}{[A]_0}$$

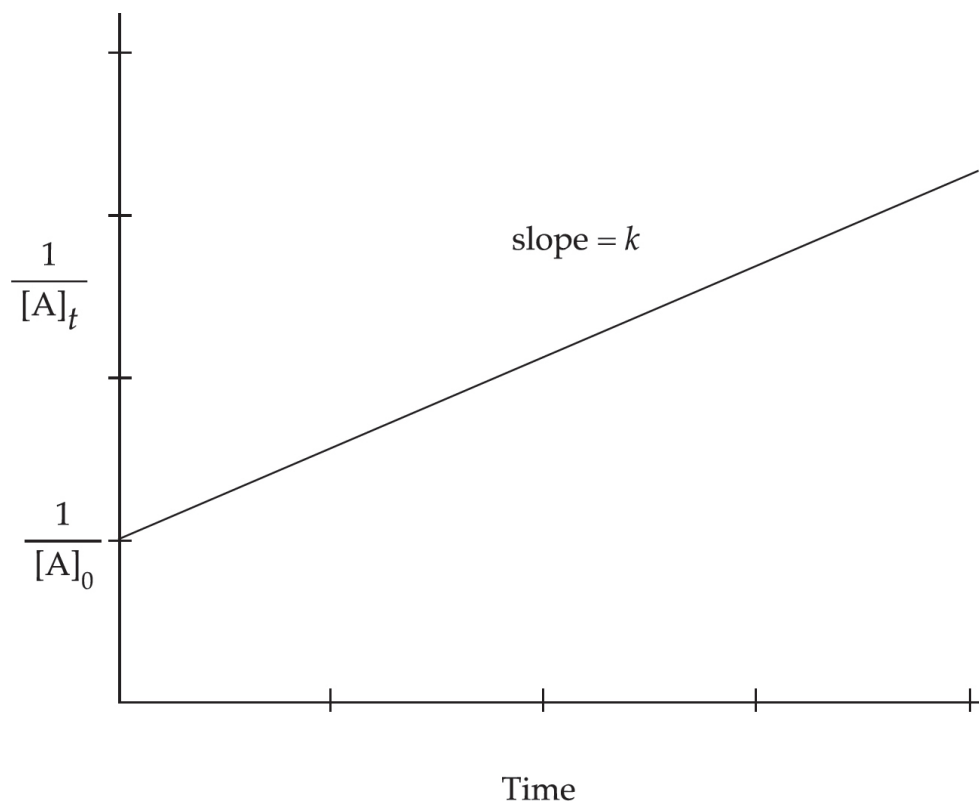
$[A]_t$ = concentration of reactant A at time t

$[A]_0$ = initial concentration of reactant A

k = rate constant

t = time elapsed

The use of inverses in the rate law creates a linear graph comparing concentration and time. Notice that the line moves upward as the concentration decreases. The slope of the line is given by k and the y -intercept is given by $\frac{1}{[A]_0}$.



Half-Life

The half-life of a reactant in a chemical reaction is the time it takes for half of the substance to react. Most half-life-problems can be solved by using a simple chart:

Time	Sample
0	100%
1 half-life	50%
2 half-lives	25%
3 half-lives	12.5%

**Reminder**

Don't forget that the chart should start with the time at zero.

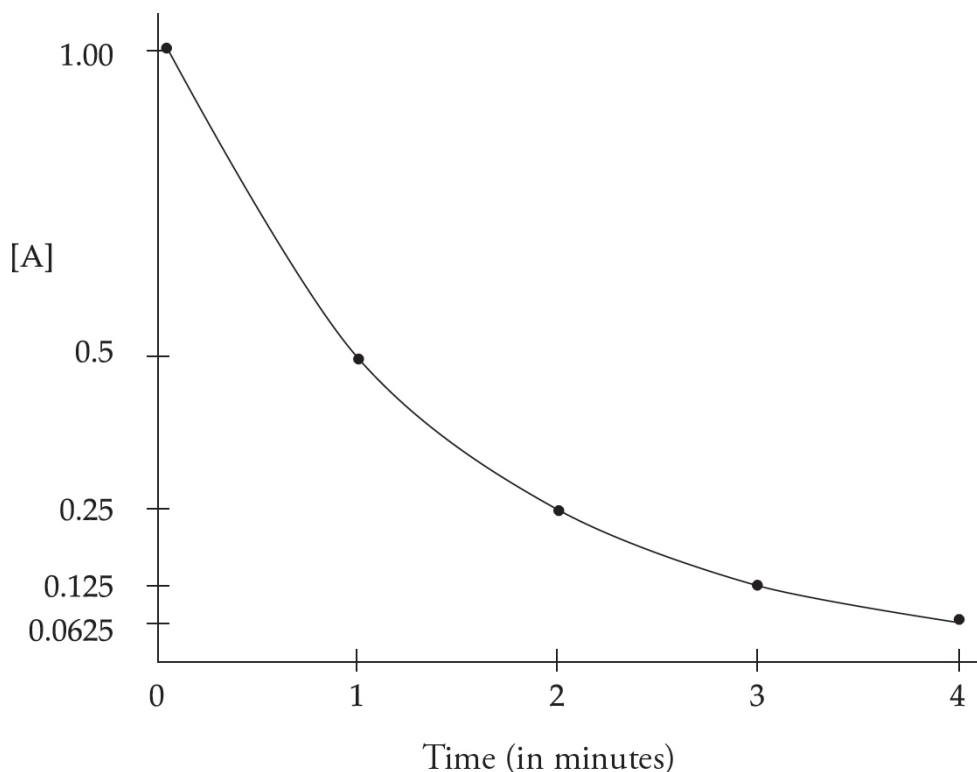
So a sample with a mass of 120 grams and a half-life of 3 years will decay as follows:

Time	Sample
0	120 g
3 years	60 g
6 years	30 g
9 years	15 g

For a first order reactant, half-life remains constant. That means if it takes 30 seconds for 50% of a reactant to decay (from 100% to 50%), then in the next 30 seconds 50% more will decay, leaving 25% of the original amount remaining. Thirty seconds later, only 12.5% will remain, etc.

This can be demonstrated graphically as shown below:

$$\text{Half-life} = \frac{\ln 2}{k} = \frac{0.693}{k}$$



Note that for zero order or second order reactants, half-life is NOT constant. You cannot use the half-life equation for anything other than a first order reactant. Zero order and second order reactants have half-lives that vary over the course of the reaction. In other words, the time it takes for the amount of a zero or second order reactant to decrease from 100% to 50% will NOT be the same as it takes for the reactant to decrease from 50% to 25%.

Let's try an example based on the data below.

[A] (M)	Time (min)
2.0	0
1.6	5
1.2	10

(a) Let's find the value of k . We'll use the first two lines of the table.

$$\ln[A]_t - \ln[A]_0 = -kt$$

$$\ln(1.6) - \ln(2.0) = -k(5 \text{ min})$$

$$-0.22 = -(5 \text{ min})k$$

$$k = 0.045 \text{ min}^{-1}$$

(b) Now let's use k to find $[A]$ when 20 minutes have elapsed.

$$\ln[A]_t - \ln[A]_0 = -kt$$

$$\ln[A]_t - \ln(2.0) = -(0.045 \text{ min}^{-1})(20 \text{ min})$$

$$\ln[A]_t = -0.21$$

$$[A]_t = e^{-0.21} = 0.81 M$$

(c) Now let's find the half-life of the reaction.

We can look at the answer (b) to see that the concentration dropped by half (1.6 M to 0.8 M) from 5 minutes to 20 minutes. That makes the half-life about 15 minutes. We can confirm this using the half-life equation.

$$\text{Half-life} = \frac{0.693}{k} = \frac{0.693}{0.045 \text{ min}^{-1}} = 15.4 \text{ minutes}$$

Another use of half-life is to examine the rate of decay of a radioactive substance. A radioactive substance is one that will slowly decay into a more stable form as time goes on.

COLLISION THEORY

According to collision theory, chemical reactions occur because reactants are constantly moving around and colliding with one another.

When reactants collide with sufficient energy (**activation energy**, E_a), a reaction occurs. These collisions are referred to as effective collisions, because they lead to a chemical reaction. Ineffective collisions do not produce a chemical reaction. At any given time

during a reaction, a certain fraction of the reactant molecules will collide with sufficient energy to cause a reaction between them.

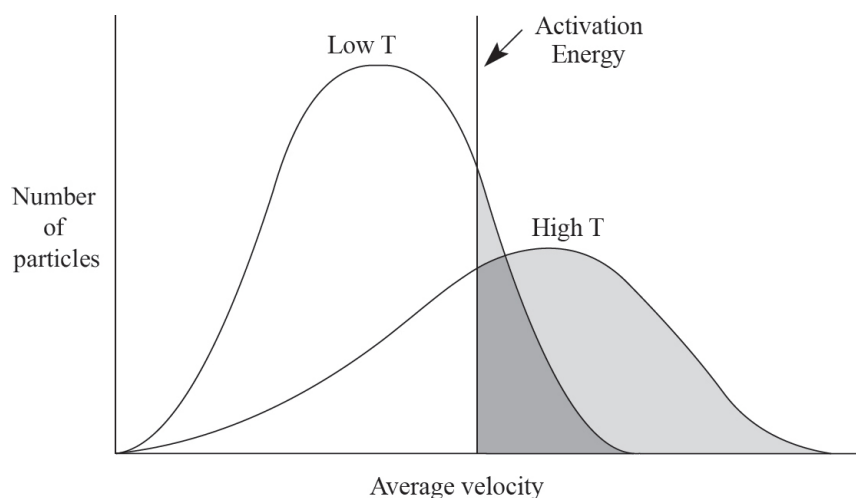
For gaseous or aqueous reactants, an increased concentration will increase the rate of reaction. This is because with more molecules moving around in a given volume, they are more likely to collide. If one of the reactants is a solid, the reaction will proceed faster if the surface area available to react is larger. For example, dropping a large chunk of metal into acid will cause the metal to dissolve, but slowly, as the acid can only react with the metal atoms on the surface. If you were to grind the metal into a powder before dropping it into the acid, the acid can react with a lot more metal atoms, and the metal would dissolve faster.

One other physical factor that can affect a reaction rate is stirring. It's certainly true that if you were to drop sugar into water, stirring the water will make the sugar dissolve faster. To explore why stirring a mixture doesn't always speed up a reaction, though, we need to talk about the concept of homogeneity. A **heterogeneous mixture** is one in which all parts of a mixture would not be identical. Before the sugar dissolves, some portions of the water will contain more sugar molecules than others, and the mixture would be heterogeneous. After the sugar fully dissolves, though, each portion of water will contain the same number of aqueous sugar molecules. A mixture in which all portions are identical is called **homogeneous**.

Stirring only increases the reaction rate in a heterogeneous mixture. Causing the solid sugar molecules to move around will make them collide with the liquid water molecules more often, which increases the likelihood that a reaction will occur. However, after the sugar dissolves completely and the mixture is homogenous, any additional sugar/water molecule collisions caused by stirring it are negligible compared to the number of collisions that are already happening due to the inherent motion of the molecules in their aqueous state.

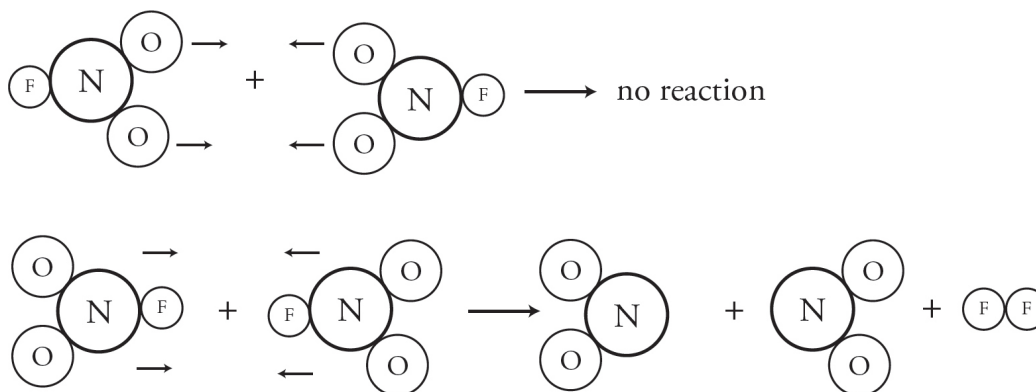
Reaction rate increases with increasing temperature because increasing temperature means that the molecules are moving faster, which means that the molecules have greater average kinetic energy. The higher the temperature, the greater the number of reactant molecules colliding with each other with enough energy (E_a) to cause a reaction.

This can be demonstrated visually on a Maxwell-Boltzmann diagram.



On the above graph, you can see that for the reaction at the higher temperature, a larger fraction of the reactant molecules have sufficient energy to exceed the activation energy barrier.

In addition, reactions only occur if the reactants collide with the correct orientation. For example, in the reaction $2\text{NO}_2\text{F} \rightarrow 2\text{NO}_2 + \text{F}_2$, there are many possible collision orientations. Two of them are drawn below:



The reaction will only occur if the two NO_2F molecules collide in such a fashion that the N–F bonds can break and the F–F bonds can form. While there is no way to quantify collision orientation, it is an important part of collision theory, and you should be familiar with the basic concept underlying it.

BEER'S LAW

To measure the concentration of a solution over time, a device called a spectrophotometer can be used in some situations. A spectrophotometer measures the amount of light at a given wavelength that is absorbed by a solution. If a solution changes color as the reaction progresses, the amount of light that is absorbed will change. Absorbance can be calculated using Beer's Law:

Beer's Law

$$A = abc$$

A = absorbance

a = molar absorptivity, a constant that depends on the solution

b = path length, the distance the light is traveling through the solution

c = concentration of the solution

Please Note

It is illegal for you to use Beer's Law until you turn 21. Only kidding! You can use Beer's Law anytime you like. It pairs well with solutions that visibly change color. (We can't stop with the jokes.)

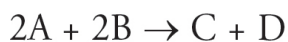
As molar absorptivity and path length are constants when using a spectrophotometer, Beer's Law is often interpreted as a direct relationship between absorbance and the concentration of the solution. Beer's Law is most effective with solutions that visibly change color over the course of a reaction, but if a spectrophotometer that emits light in the ultraviolet region is used, Beer's Law can be used to determine the concentrations of reactants in solutions that are invisible to the human eye.

You may also run into a device called a colorimeter while studying Beer's Law. A colorimeter is simply a spectrophotometer that can only emit light at specific frequencies, whereas a spectrophotometer can emit light at any frequency within a set range.

REACTION MECHANISMS

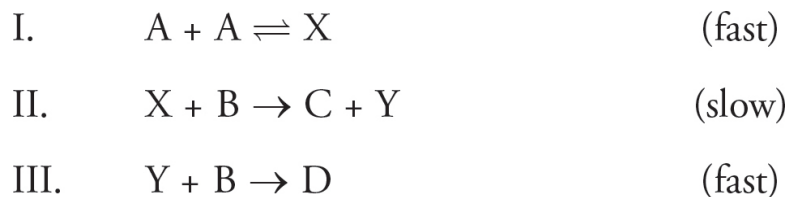
Many chemical reactions are not one-step processes. Rather, the balanced equation is the sum of a series of individual steps, called elementary steps.

For instance, the hypothetical reaction

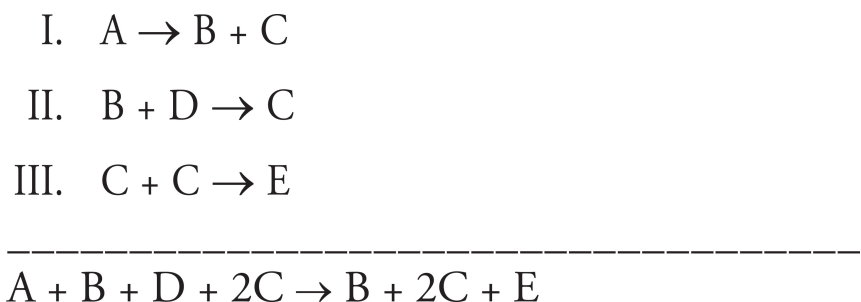


$$\text{Rate} = k[A]^2[B]$$

could take place by the following three-step mechanism:



Species X and Y are called reaction **intermediates**, because they are produced but also fully consumed over the course of the reaction. Intermediates will always cancel out when adding up the various elementary steps in a reaction, as shown below.



Cancel the species that appear on both sides.

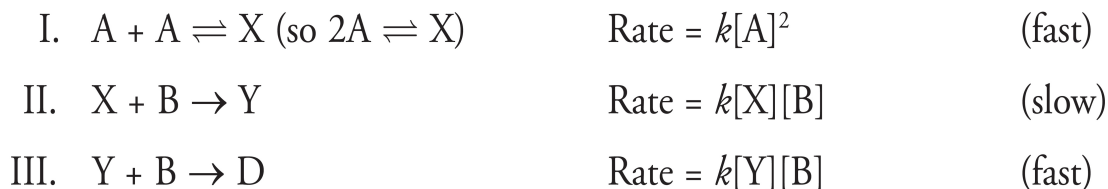


By adding up all the steps, we get the balanced equation for the overall reaction, so this mechanism is consistent with the balanced equation. All elementary steps have either one or two reactants. Any elementary step that has two reactants (even if they are the same!) is called **bimolecular**. Any elementary step that only has one reactant is called **unimolecular**. In the above mechanism, step I is unimolecular and steps II and III are bimolecular.

As in any process where many steps are involved, the speed of the whole process can't be faster than the speed of the slowest step in the process, so the slowest step of a reaction is called the **rate-determining step**. Since the slowest step is the most important step in determining the rate of a reaction, the slowest step and the

steps leading up to it are used to see if the mechanism is consistent with the rate law for the overall reaction.

The rate for an elementary step can be determined by taking the concentration of the reactants in that step and raising them to the power of any coefficient attached to that reactant. Let's look at an example:

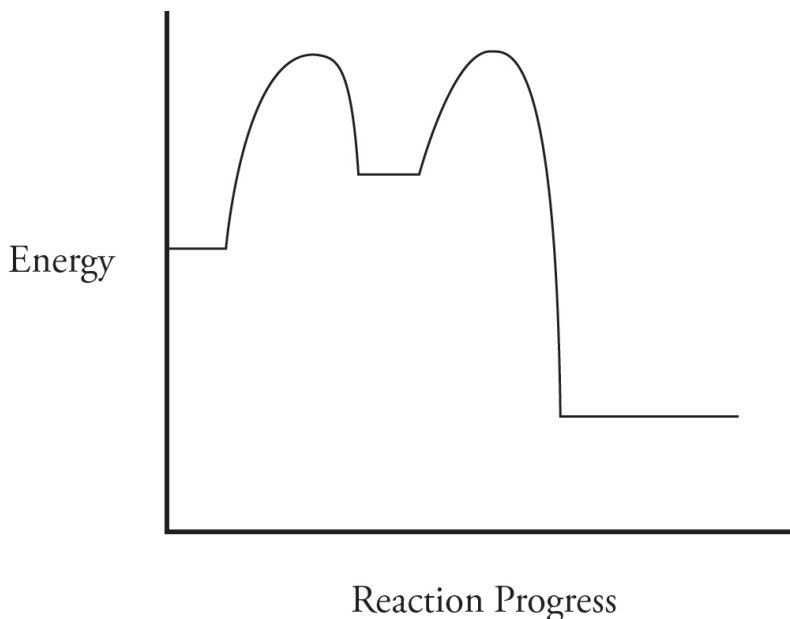


The rate law for the entire reaction is equal to that of the slowest elementary step, which is step II. However, step II has an intermediate (X) present in it. Even though intermediates can appear in rate laws, if they can be replaced with a reactant via an earlier step, we should try and do so. Looking at step 1, we can also see that [X] is equivalent to $[A]^2$ (as the sides are in equilibrium). If $[A]^2$ is substituted in for [X], the rate law for step II becomes rate = $k[A]^2[B]$, which is the overall rate law for this reaction.

It is important to emphasize that you can only use the coefficient method to determine the rate law of elementary steps. You cannot use the coefficients from the overall balanced equation in a similar fashion to determine the overall rate law. (For instance, the rate law for the above example is not rate = $k[A]^2[B]^2$). The only way to use the coefficient method to determine the rate law for a full reaction is by knowing which elementary step is the slowest and applying the above method to that step. If you do not know the relative speed of the elementary steps, you must use experimental data to determine the rate law.

As discussed previously, an energy diagram is a graphical representation that shows the energy level of the products and the reactants, as well as the required activation energy for a reaction to

occur. As many reactions take place over several steps, an energy diagram can be broken down to examine the energy change in each individual step. Take the diagram below:



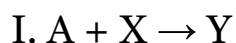
In this diagram, you can see the overall reaction is exothermic. However, the reaction takes place in two different steps: the first, which is endothermic, and the second, which is exothermic.

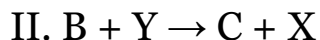
CATALYSTS

As we mentioned earlier in the book, a catalyst increases the rate of a chemical reaction without being consumed in the process; catalysts do not appear in the balanced equation. In some cases, a catalyst is a necessary part of a reaction because in its absence, the reaction would proceed at too slow of a rate to be at all useful.

When looking at elementary steps, a catalyst is present both before and after the overall reaction. Catalysts cancel out of the overall reaction but are present in elementary steps.

Example: $A + B \rightarrow C$

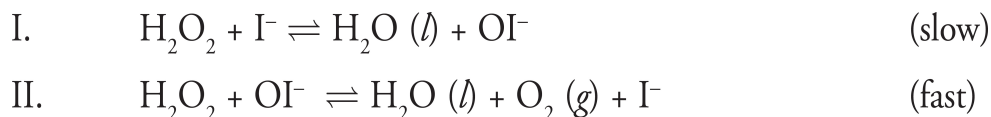




In the above example, X is a catalyst and Y is a reaction intermediate.

A catalyst increases the rate of a chemical reaction by providing an alternative reaction pathway with a lower activation energy.

Note that catalysts and (very rarely) intermediates can appear in rate laws. Let's look at the decomposition of H_2O_2 in the presence of the iodide ion as an example. The two elementary steps are shown below:



If you combine those steps, the overall reaction becomes $\text{H}_2\text{O}_2 (aq) \rightleftharpoons 2 \text{H}_2\text{O} (l) + \text{O}_2 (g)$. In the given mechanism, I^- is a catalyst, and OI^- is an intermediate. As I^- appears as a reactant in the slow step, it would be part of the overall rate law, which would be $\text{rate} = k[\text{H}_2\text{O}_2][\text{I}^-]$.

When a catalyst is introduced to a reaction, the ensuing reaction is said to undergo catalysis. There are many types of catalysis. One of the most common is surface catalysis, in which a reaction intermediate is formed as in the example above. Another is enzyme catalysis, in which the catalyst binds to the reactants in a way to reduce the overall activation energy of the reaction. Enzymes are very common in biological applications. Finally, in acid-base catalysis, reactants will gain or lose protons in order to change the rate of reaction. Acids and bases will be studied in more depth in Big Idea #6.

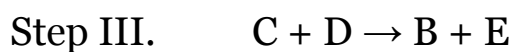
CHAPTER 6 QUESTIONS

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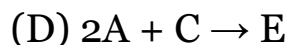
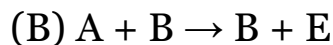
Multiple-Choice Questions

Use the following information to answer questions 1-4.

A multi-step reaction takes place with the following elementary steps:



1. What is the overall balanced equation for this reaction?



2. What is the function of species B in this reaction?

(A) Without it, no reaction would take place.

(B) It is a reaction intermediate which facilitates the progress of the reaction.

(C) It is a catalyst which changes the overall order of the reaction.

(D) It lowers the overall activation energy of the reaction.

3. If step II is the slow step for the reaction, what is the overall rate law?

(A) $\text{Rate} = k[\text{A}]^2[\text{B}]$

(B) $\text{Rate} = k[\text{A}][\text{C}]$

(C) $\text{Rate} = k[\text{A}][\text{B}]$

(D) $\text{Rate} = k[\text{A}]/[\text{D}]$

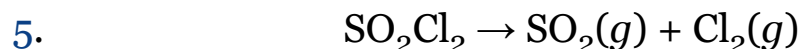
4. Why would increasing the temperature make the reaction rate go up?

(A) It is an endothermic reaction that needs an outside energy source to function.

(B) The various molecules in the reactions will move faster and collide more often.

(C) The overall activation energy of the reaction will be lowered.

(D) A higher fraction of molecules will have the same activation energy.



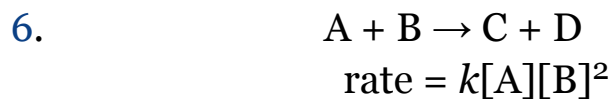
At 600 K, SO_2Cl_2 will decompose to form sulfur dioxide and chlorine gas via the above equation. If the reaction is found to be first order overall, which of the following will cause an increase in the half life of SO_2Cl_2 ?

(A) Increasing the initial concentration of SO_2Cl_2

(B) Increasing the temperature at which the reaction occurs

(C) Decreasing the overall pressure in the container

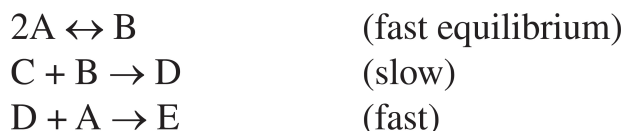
(D) None of these will increase the half life.



What are the potential units for the rate constant for the above reaction?

- (A) s^{-1}
- (B) $s^{-1}M^{-1}$
- (C) $s^{-1}M^{-2}$
- (D) $s^{-1}M^{-3}$

7. The following mechanism is proposed for a reaction:



Which of the following is the correct rate law for complete reaction?

- (A) $\text{Rate} = k[C]^2[B]$
- (B) $\text{Rate} = k[C][A]^2$
- (C) $\text{Rate} = k[C][A]^3$
- (D) $\text{Rate} = k[D][A]$



The reaction above takes place with all of the reactants and products in the gaseous phase. Which of the following is true of the relative rates of disappearance of the reactants and appearance of the products?

- (A) NO appears at twice the rate that NOCl disappears.
- (B) NO appears at the same rate that NOCl disappears.
- (C) NO appears at half the rate that NOCl disappears.
- (D) Cl_2 appears at the same rate that NOCl disappears.

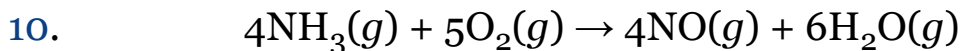


When the reaction given above takes place in a sealed isothermal container, the rate law is

$$\text{Rate} = k[\text{H}_2][\text{I}_2]$$

If a mole of H_2 gas is added to the reaction chamber and the temperature remains constant, which of the following will be true?

- (A) The rate of reaction and the rate constant will increase.
- (B) The rate of reaction and the rate constant will not change.
- (C) The rate of reaction will increase and the rate constant will decrease.
- (D) The rate of reaction will increase and the rate constant will not change.



The above reaction will experience a rate increase by the addition of a catalyst such as platinum. Which of the following best explains why?

- (A) The catalyst causes the value for ΔG to become more negative.
- (B) The catalyst increases the percentage of collisions that occur at the proper orientation in the reactant molecules.
- (C) The catalyst introduces a new reaction mechanism for the reaction.
- (D) The catalyst increases the activation energy for the reaction.



Based on the following experimental data, what is the rate law for the hypothetical reaction given above?

Experiment	[A] (M)	[B] (M)	Initial Rate of Formation of C (mol/L·sec)
1	0.20	0.10	3×10^{-2}
2	0.20	0.20	6×10^{-2}
3	0.40	0.20	6×10^{-2}

- (A) $\text{Rate} = k[\text{A}]$
- (B) $\text{Rate} = k[\text{A}]^2$
- (C) $\text{Rate} = k[\text{B}]$
- (D) $\text{Rate} = k[\text{A}][\text{B}]$

12. A solution of Co^{2+} ions appears red when viewed under white light. Which of the following statements is true about the solution?

- (A) A spectrophotometer set to the wavelength of red light would read a high absorbance.
- (B) If the solution is diluted, the amount of light reflected by the solution will decrease.
- (C) All light with a frequency that is lower than that of red light will be absorbed by it.
- (D) Electronic transmissions within the solution match the wavelength of red light.

13.



Based on the following experimental data, what is the rate law for the hypothetical reaction given above?

Experiment	[A] (<i>M</i>)	[B] (<i>M</i>)	Initial Rate of Formation of C (<i>M</i> /sec)
1	0.20	0.10	2.0×10^{-6}
2	0.20	0.20	4.0×10^{-6}
3	0.40	0.40	1.6×10^{-5}

(A) Rate = $k[A]$

(B) Rate = $k[A]^2$

(C) Rate = $k[B]$

(D) Rate = $k[A][B]$

14.

Time (Hours)	[A] <i>M</i>
0	0.40
1	0.20
2	0.10
3	0.05

Reactant A underwent a decomposition reaction. The concentration of A was measured periodically and recorded in the chart above. Based on the data in the chart, which of the following is the rate law for the reaction?

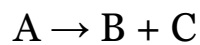
(A) Rate = $k[A]$

(B) $\text{Rate} = k[\text{A}]^2$

(C) $\text{Rate} = 2k[\text{A}]$

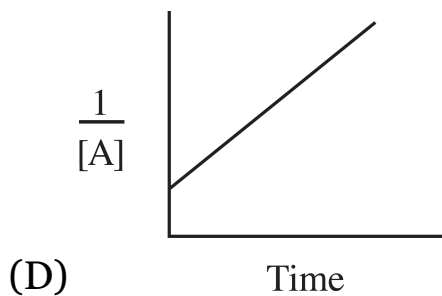
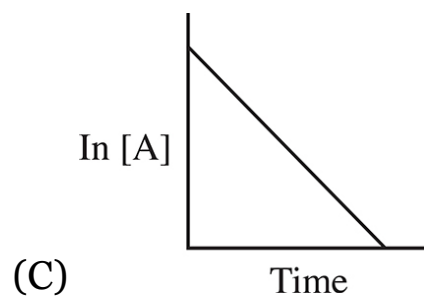
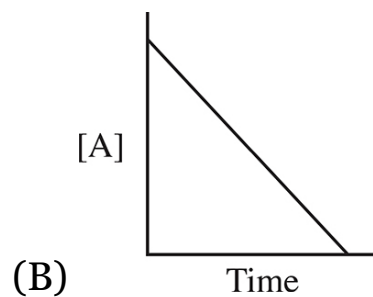
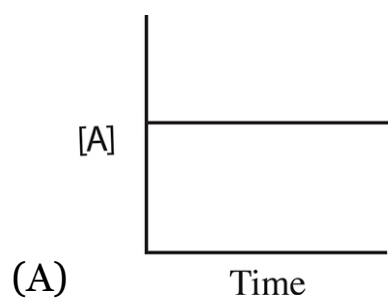
(D) $\text{Rate} = \frac{1}{2}k[\text{A}]$

15.



$$\text{rate} = k[\text{A}]^2$$

Which of the following graphs may have been created using data gathered from the above reaction?



16. After 44 minutes, a sample of $^{44}_{19}\text{K}$ is found to have decayed to 25 percent of the original amount present. What is the half-life of $^{44}_{19}\text{K}$?

- (A) 11 minutes
(B) 22 minutes
(C) 44 minutes
(D) 66 minutes

Free-Response Questions

1. $\text{A} + 2\text{B} \rightarrow 2\text{C}$

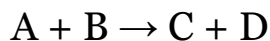
The following results were obtained in experiments designed to study the rate of the reaction above:

Experiment	Initial Concentration (mol/L)		Initial Rate of Disappearance of A (M/sec)
	[A]	[B]	
1	0.05	0.05	3.0×10^{-3}
2	0.05	0.10	6.0×10^{-3}
3	0.10	0.10	1.2×10^{-2}
4	0.20	0.10	2.4×10^{-2}

- (a) Determine the order of the reaction with respect to each of the reactants, and write the rate law for the reaction.
- (b) Calculate the value of the rate constant, k , for the reaction. Include the units.
- (c) If another experiment is attempted with [A] and [B], both 0.02-molar, what would be the initial rate of disappearance of

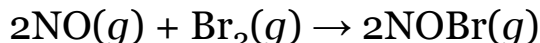
A?

- (d) The following reaction mechanism was proposed for the reaction above:



- (i) Show that the mechanism is consistent with the balanced reaction.
- (ii) Show which step is the rate-determining step, and explain your choice.

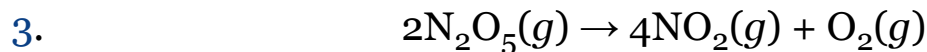
2.



The following results were obtained in experiments designed to study the rate of the reaction above:

Experiment	Initial Concentration (mol/L)		Initial Rate of Appearance of NOBr (M/sec)
	[NO]	[Br ₂]	
1	0.02	0.02	9.6×10^{-2}
2	0.04	0.02	3.8×10^{-1}
3	0.02	0.04	1.9×10^{-1}

- (a) Write the rate law for the reaction.
- (b) Calculate the value of the rate constant, k , for the reaction. Include the units.
- (c) In experiment 2, what was the concentration of NO remaining when half of the original amount of Br₂ was consumed?
- (d) Which of the following reaction mechanisms is consistent with the rate law established in (a)? Explain your choice.

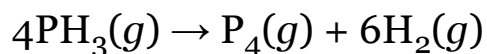


Dinitrogen pentoxide gas decomposes according to the equation above. The first-order reaction was allowed to proceed at 40°C and the data below were collected.

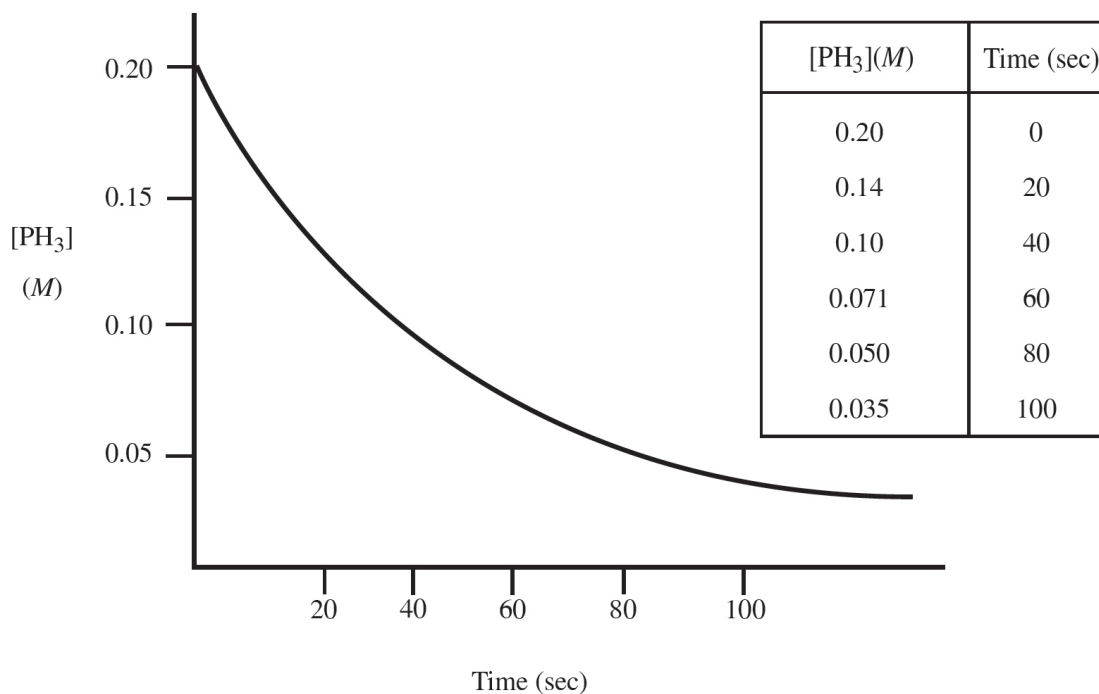
$[\text{N}_2\text{O}_5]$ (M)	Time (min)
0.400	0.0
0.289	20.0
0.209	40.0
0.151	60.0
0.109	80.0

- Calculate the rate constant for the reaction using the values for concentration and time given in the table. Include units with your answer.
- After how many minutes will $[\text{N}_2\text{O}_5]$ be equal to 0.350 M ?
- What will be the concentration of N_2O_3 after 100 minutes have elapsed?
- Calculate the initial rate of the reaction. Include units with your answer.
- What is the half-life of the reaction?

4. The decomposition of phosphine occurs via the pathway below:



A scientist observing this reaction at 250 K plots the following data:



- (a) (i) What order is this reaction? Why?
- (ii) What would the concentration of the PH_3 gas be after 120 sec?
- (b) If the rate of disappearance of PH_3 is $2.5 \times 10^{-3} \text{ M/s}$ at $t = 20$ s:
- (i) What is the rate of appearance of P_4 at the same point in time?
- (ii) How will the rate of appearance of P_4 change as the reaction progresses forward?
- (c) The experiment is repeated with the same initial concentration of phosphine, but this time the temperature is set at 500 K. How and why would the following values change, if at all?

- (i) The half-life of the phosphine
- (ii) The rate law
- (iii) The value of the rate constant

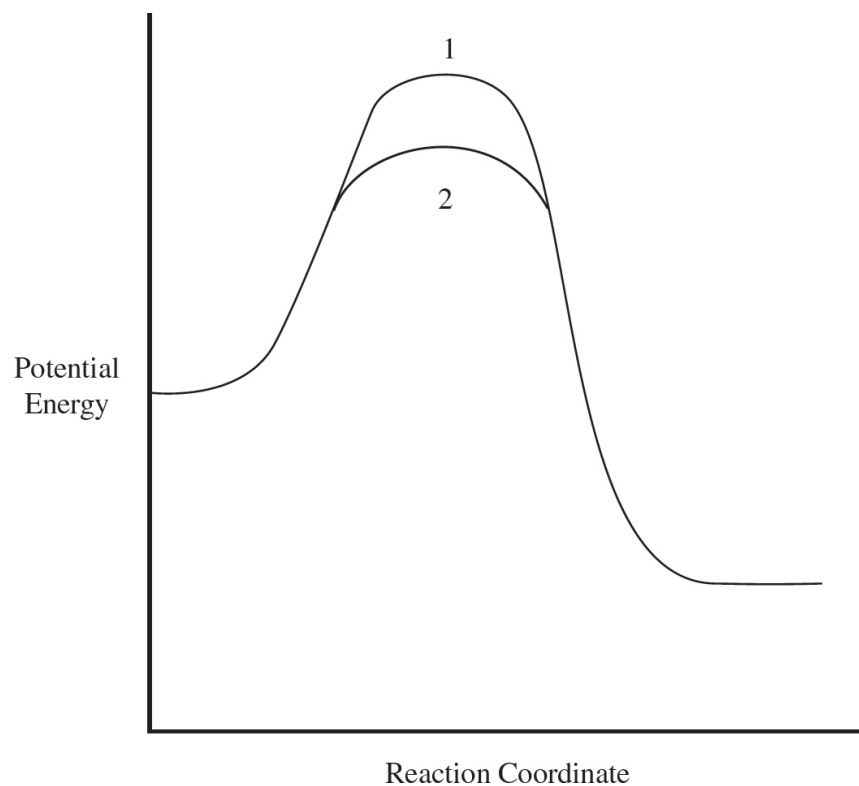


The reaction above is second order with respect to A and zero order with respect to B. Reactants A and B are present in a closed container. Predict how each of the following changes to the reaction system will affect the rate and rate constant, and explain why.

- (a) More gas A is added to the container.
- (b) More gas B is added to the container.
- (c) The temperature is increased.
- (d) An inert gas D is added to the container.
- (e) The volume of the container is decreased.

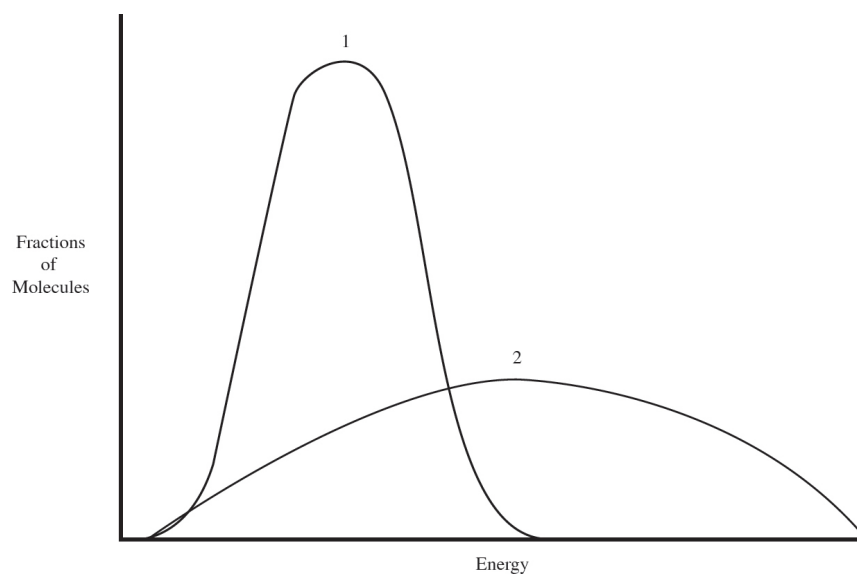
6. Use your knowledge of kinetics to answer the following questions. Justify your answers.

- (a)



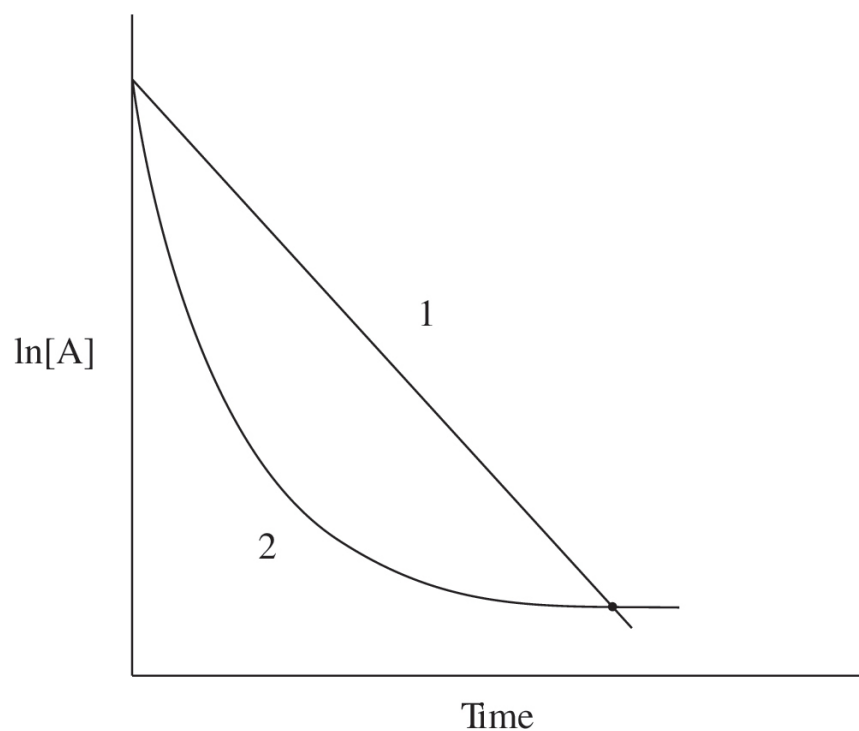
The two lines in the diagram above show different reaction pathways for the same reaction. Which of the two lines shows the reaction when a catalyst has been added?

(b)



Which of the two lines in the energy distribution diagram shows the conditions at a higher temperature?

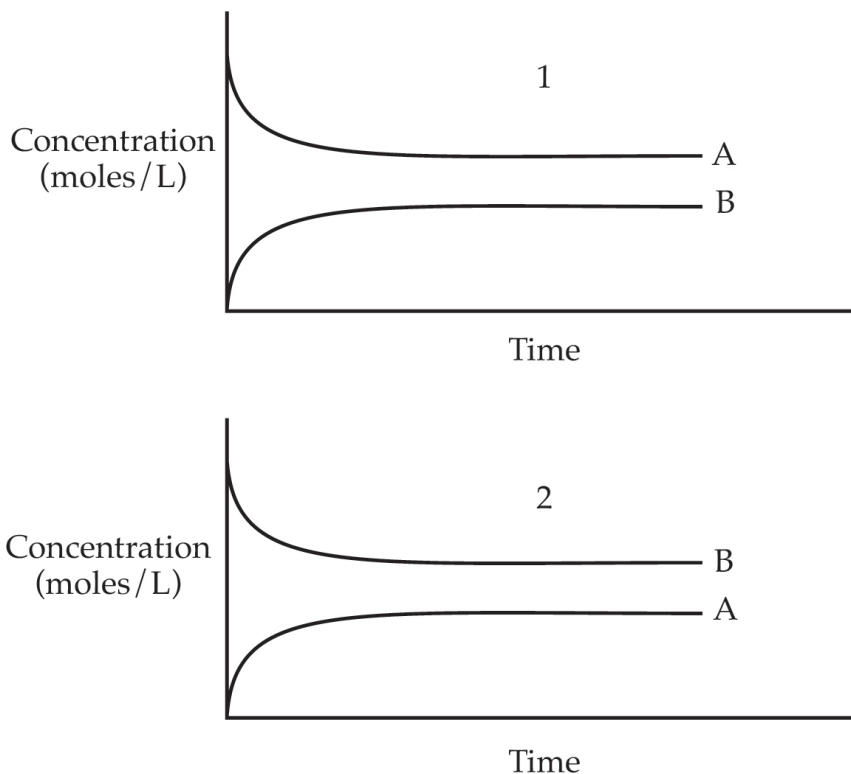
(c)



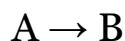
Which of the two lines in the diagram above shows the relationship of $\ln[A]$ to time for a first order reaction with the following rate law?

$$\text{Rate} = k[A]$$

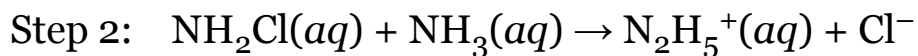
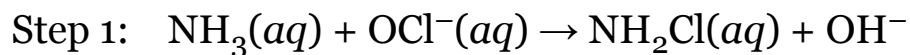
(d)



Which of the two graphs above shows the changes in concentration over time for the following reaction?



7. Hydrazine (N_2H_4) can be produced commercially via the Raschig process. The following is a proposed mechanism:



- (a) (i) What is the equation for the overall reaction?
 (ii) Identify any catalysts or intermediates from the reaction mechanism.
- (b) The rate law for the reaction is determined to be $\text{rate} = k[NH_3][OCl^-]$

(i) Which elementary step is the slowest one? Justify your answer.

(ii) If the reaction is measured over the course of several minutes, what would the units of the rate constant be?

8. The reaction between crystal violet (a complex organic molecule represented by CV^+) and sodium hydroxide is as follows:



As the crystal violet is the only colored species in the reaction, a spectrophotometer calibrated to a specific wavelength can be used to determine its concentration over time. The following data was gathered:

$[\text{CV}^+] \text{ (M)}$	Time (s)
5.5×10^{-5}	0
3.8×10^{-5}	60
2.6×10^{-5}	120
1.8×10^{-5}	180

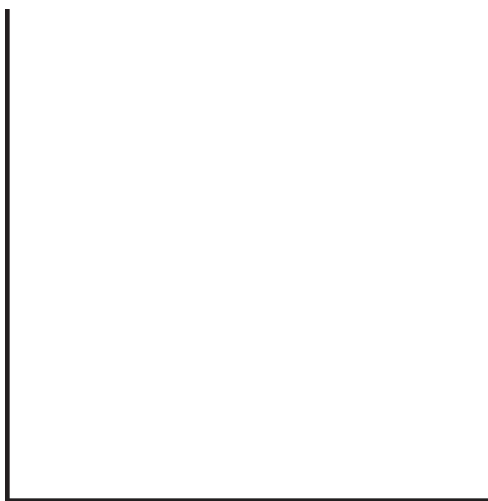
(a) (i) What is the rate of disappearance for crystal violet from $t = 60 \text{ s}$ to $t = 120 \text{ s}$?

(ii) If the solution placed in the spectrophotometer 30 s after mixing instead of immediately after mixing, how would that affect the calculated rate of disappearance for crystal violet in part (i)?

(b) Given the path length of the cuvette is 1.00 cm and the molar absorptivity of the solution is $26,000 \text{ cm}^{-1}\text{M}^{-1}$ at the

wavelength of the spectrophotometer, what would the absorbance reading on the spectrophotometer be at $t = 60$ s?

- (c) This reaction is known to be first order with respect to crystal violet. On the provided axes, graph a function of $[\text{CV}^+]$ vs. time that will provide you with a straight line graph.



The following data was also gathered over the course of three experiments:

Experiment	$[\text{CV}^+]_{\text{init}}$ (M)	$[\text{OH}^-]_{\text{init}}$ (M)	Initial rate of formation of CVOH (M/s)
1	5.5×10^{-5}	0.12	3.60×10^{-7}
2	5.5×10^{-5}	0.24	7.20×10^{-7}
3	4.1×10^{-5}	0.18	?

- (d) Write the rate law for this reaction.
- (e) What is the rate constant, k , for this reaction? Include units in your answer.
- (f) Determine the initial rate of formation of CVOH for experiment 3.

CHAPTER 6 ANSWERS AND EXPLANATIONS

Multiple-Choice

1. **D**

The answer to (A) shows all of the reactants and products that are part of this reaction. However, species that appear on both sides need to cancel. That eliminates B, C, and D, leaving $2A + C \rightarrow E$

2. **D**

Species B is a catalyst, as it is present before and after the reaction but cancels out of the overall equation. However, catalysts do not change the order of the reaction; they simply lower the activation energy so the reaction can proceed more quickly. The reaction would still take place without the catalyst, but it would proceed very slowly.

3. **A**

Only reactants can appear in rate laws. In Step II, you can replace C with A + B, as these are shown to be equivalent in Step I. Step II now reads:



As Step II is the slow step, the coefficients then become exponents in the overall rate law, as in (A).

4. **B**

Increasing the temperature increases the speed of the molecules. This will increase how often those molecules collide, meaning that a successful collision is more likely to

happen. Additionally, there will also be more energy within each collision, making it more likely that the collision will exceed the activation energy barrier.

5. **D**

None of the options would decrease the rate of reaction, which would be required for the half-life of the reactant to increase.

6. **C**

To answer this, you can do a unitless dimensional analysis:

$$M/s = k(M)(M)^2$$

$$M/s = k(M)^3$$

$$k = s^{-1}M^{-2}$$

7. **B**

The coefficients on the reactants in the slowest elementary step match up with the order of those reactants in the overall rate law. However, B is an intermediate that is not present at the start of the reaction and as such cannot be part of the rate law. B is in equilibrium with 2A in the first step, though, so the 2A can be substituted for B in the slow step, which then yields (B) when the rate law is determined.

8. **B**

For every two NO molecules that form, two NOCl molecules must disappear, so NO is appearing at the same rate that NOCl is disappearing. Choice (D) is wrong because for every mole of Cl_2 that forms, two moles of

NOCl are disappearing, so Cl_2 is appearing at *half* the rate that NOCl is disappearing.

9. **D**

From the rate law given in the question ($\text{Rate} = k[\text{H}_2][\text{I}_2]$), you can see that increasing the concentration of H_2 will increase the rate of reaction. The rate constant, k , is not affected by changes in the concentration of the reactants.

10. **C**

Catalysts work by creating a new reaction pathway with a lower activation energy than the original pathway.

11. **C**

From a comparison of experiments 1 and 2, when $[\text{B}]$ is doubled while $[\text{A}]$ is held constant, the rate doubles. That means that the reaction is first order with respect to B.

From a comparison of experiments 2 and 3, when $[\text{A}]$ is doubled while $[\text{B}]$ is held constant, the rate doesn't change. That means that the reaction is zero order with respect to A and that A will not appear in the rate law.

So the rate law is $\text{Rate} = k[\text{B}]$.

12. **D**

A spectrophotometer works by reading light absorbance. A red solution appears to be red because it transmits red light. The amount of light reflected by the solution does not change as it dilutes; the amount of light that is transmitted and absorbed does. The solution has a red color because the distance between the energy levels within the cobalt ions corresponds with the wavelength of red light.

13. **D**

From a comparison of experiments 1 and 2, when [B] is doubled while [A] is held constant, the rate doubles. That means that the reaction is first order with respect to B.

From a comparison of experiments 2 and 3, when both [A] and [B] are doubled, the rate increases by a factor of 4. We would expect the rate to double based on the change in B; because the rate is in fact multiplied by 4, the doubling of A must also change the rate by a factor of 2, so the reaction is also first order with respect to A.

So the rate law is $\text{Rate} = k[\text{A}][\text{B}]$.

14. **A**

The key to this question is to recognize that reactant A is disappearing with a characteristic half-life. This is a signal that the reaction is first order with respect to A. So the rate law must be $\text{Rate} = k[\text{A}]$.

15. **D**

The reaction is a second order reaction, of which a graph of the inverse of concentration always produces a straight line with a slope equal to k , the rate constant.

16. **B**

Make a chart. Always start at time = 0.

Half-Lives	Time	Stuff
0	0	100%
1	X	50%

2	44 min.	25%
---	---------	-----

It takes two half-lives for the amount of to decrease to 25 percent. If two half-lives takes 44 minutes, one half-life must be 22 minutes.

Free-Response

- (a) Comparing the results of experiments 3 and 4, you can see that when $[A]$ doubles, the rate doubles, so the reaction is first order with respect to A.

When you compare the results of experiments 1 and 2, you can see that when $[B]$ doubles, the rate doubles, so the reaction is first order with respect to B.

$$\text{Rate} = k[A][B]$$

- (b) Use the values from experiment 3, just because they look the simplest.

$$k = \frac{\text{Rate}}{[A][B]} = \frac{(1.2 \times 10^{-2} \text{ M/sec})}{(0.10 \text{ M})(0.10 \text{ M})} = 1.2 \text{ M}^{-1}\text{sec}^{-1}$$

- (c) Use the rate law.

$$\text{Rate} = k[A][B]$$

$$\text{Rate} = (1.2 \text{ M}^{-1}\text{sec}^{-1})(0.02 \text{ M})(0.02 \text{ M}) = 4.8 \times 10^{-4} \text{ M/sec}$$

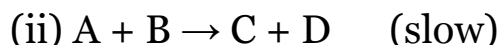
- (d) (i) $A + B \rightarrow C + D$



The two reactions add up to



D's cancel, and you're left with the balanced equation.



The first part of the mechanism is the slow, rate-determining step because its rate law is the same as the experimentally determined rate law.

2. (a) Comparing the results of experiments 1 and 2, you can see that when $[NO]$ doubles, the rate quadruples, so the reaction is second order with respect to NO .

Comparing the results of experiments 1 and 3, you can see that when $[Br_2]$ doubles, the rate doubles, so the reaction is first order with respect to Br_2 .

$$\text{Rate} = k[NO]^2[Br_2]$$

- (b) Using the values from experiment 1, we first need to realize that the rate of reaction would be half of the rate of appearance of $NOBr$, because of the "2" coefficient in front of the $NOBr$. So, the rate of reaction for experiment 1 would be $0.0096/2 = 0.0048 \text{ M/s}$. Plugging that into the rate law yields:

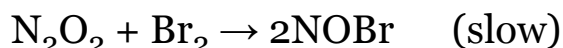
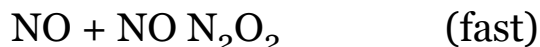
$$\text{Rate} = k[NO]^2[Br_2] \quad 0.0048 \text{ M/s} = (0.02 \text{ M})^2(0.02 \text{ M}) k = 6 \times 10^3 \text{ M}^{-2}\text{s}^{-1}$$

- (c) In experiment 2, you started with $[Br_2] = 0.02 \text{ M}$, so 0.01 M was consumed.

From the balanced equation, 2 moles of NO are consumed for every mole of Br₂ consumed. So 0.02 M of NO are consumed.

$$[\text{NO}]_{\text{remaining}} = 0.04 M - 0.02 M = 0.02 M$$

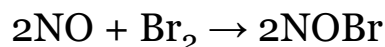
(d) Choice (I) agrees with the rate law.



The slow step is the rate-determining step, with the following rate law:

$$\text{Rate} = k[\text{N}_2\text{O}_2][\text{Br}_2]$$

Looking at the first step in the proposed mechanism, you can replace the N₂O₂ with 2NO. Doing so gives you:



The rate law for that elementary step matches the overall rate law determined previously.

3. (a) Use the First-Order Rate Law and insert the first two lines from the table.

$$\ln[\text{N}_2\text{O}_5]_t - \ln[\text{N}_2\text{O}_5]_0 = -kt$$

$$\ln(0.289) - \ln(0.400) = -k(20.0 \text{ min})$$

$$-.325 = -k(20.0 \text{ min})$$

$$k = 0.0163 \text{ min}^{-1}$$

- (b) Use the First-Order Rate Law.

$$\ln[\text{N}_2\text{O}_5]_t - \ln[\text{N}_2\text{O}_5]_0 = -kt$$

$$\ln(0.350) - \ln(0.400) = -(0.0163 \text{ min}^{-1})t$$

$$-0.134 = -(0.0163 \text{ min}^{-1})t$$

$$t = 8.22 \text{ min}$$

(c) Use the First-Order Rate Law.

$$\ln[\text{N}_2\text{O}_5] - \ln[\text{N}_2\text{O}_5]_0 = -kt$$

$$\ln[\text{N}_2\text{O}_5] - \ln(0.400) = -(0.0163 \text{ min}^{-1})(100 \text{ min})$$

$$\ln[\text{N}_2\text{O}_5] + 0.916 = 1.63$$

$$\ln[\text{N}_2\text{O}_5] = -2.55$$

$$[\text{N}_2\text{O}_5] = e^{-2.55} M = 0.078 M$$

(d) For a first-order reaction, $\text{Rate} = k[\text{N}_2\text{O}_5] = (0.0163 \text{ min}^{-1})(0.400 M) = 0.00652 M/\text{min}$.

(e) You can see from the numbers on the table that the half-life is slightly over 40 min. To calculate it exactly, use the formula

$$\text{Half-life} = \frac{0.693}{k} = \frac{0.693}{0.0163 \text{ min}^{-1}} = 42.5 \text{ min}.$$

4. (a) (i) The reaction is first order. The half-life is constant; it takes 40 seconds for the concentration to drop from 0.20 *M* to 0.10 *M*, and another 40 seconds for it to drop from 0.10 *M* to 0.050 *M*. This type of exponential decay is typical of a first order reaction.

- (ii) After $t = 80$ s, another 40 seconds is an additional half-life, and it would bring the concentration down to 0.025 M (half of 0.050 M).

- (b) (i) Use the stoichiometric ratios to determine the rate of appearance of P_4 :

$$2.5 \times 10^{-3} \text{ M/s } PH_3 \times \frac{1 \text{ mol } P_4}{4 \text{ mol } PH_3} = 6.3 \times 10^{-4} \text{ M/s } P_4$$

- (ii) As the reaction progresses, the rate at which the phosphine disappears will decrease, as indicated by the decreasing slope of the line. Thus, the rate of appearance of P_4 will also decrease.

- (c) (i) The half-life of the phosphine will decrease. This is because at a higher temperature, the reaction will proceed faster. It will thus take less time for the phosphine to decay.

- (ii) The rate law is unaffected by outside conditions and will remain unchanged.

- (iii) The rate constant will increase. There is a directly proportional relationship between the rate constant and temperature. As the temperature increases, so will the rate constant.

5. (a) The rate of the reaction will increase because the rate depends on the concentration of A as given in the rate law: $\text{Rate} = k[A]^2$.

The rate constant is independent of the concentration of the reactants and will not change.

- (b) The rate of the reaction will not change. If the reaction is zero order with respect to B, then the rate is independent of

the concentration of B.

The rate constant is independent of the concentration of the reactants and will not change.

- (c) The rate of the reaction will increase with increasing temperature because the rate constant increases with increasing temperature.

The rate constant increases with increasing temperature because at a higher temperature more gas molecules will collide with enough energy to overcome the activation energy for the reaction.

- (d) Neither the rate nor the rate constant will be affected by the addition of an inert gas.

- (e) The rate of the reaction will increase because decreasing the volume of the container will increase the concentration of A: $\text{Rate} = k[A]^2$.

The rate constant is independent of the concentration of the reactants and will not change.

6. (a) Line 2 is the catalyzed reaction. Adding a catalyst lowers the activation energy of the reaction, making it easier for the reaction to occur.
- (b) Line 2 shows the higher temperature distribution. At a higher temperature, more of the molecules will be at higher energies, causing the distribution to flatten out and shift to the right.
- (c) Line 1 is correct. The $\ln[\text{reactant}]$ for a first order reaction changes in a linear fashion over time, as shown in the following equation.

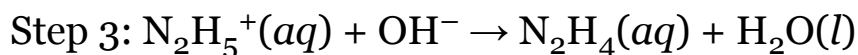
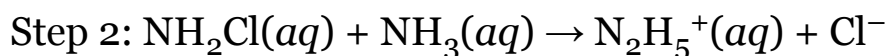
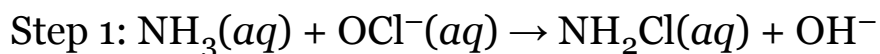
$$\ln[A]_t = -kt + \ln[A]_0$$

$$y = mx + b$$

Notice the similarity to the slope-intercept form for a linear equation.

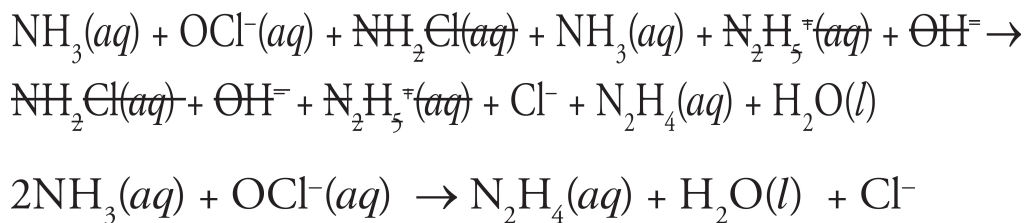
(d) Graph 1 is correct, showing a decrease in the concentration of A as it is consumed in the reaction and a corresponding increase in the concentration of B as it is produced.

7. Hydrazine (N_2H_4) can be produced commercially via the Raschig process. The following is a proposed mechanism:



(a) (i) What is the equation for the overall reaction?

To determine the net equation, all three equations must be added together, and species that appear on both sides of the arrow can be eliminated.



(ii) Identify any catalysts or intermediates from the reaction mechanism.

There are no catalysts present, but NH_2Cl , N_2H_5^+ , and OH^- are all intermediates in the process.

(b) The rate law for the reaction is determined to be $\text{rate} = k[\text{NH}_3][\text{OCl}^-]$.

(i) Which elementary step is the slowest one? Justify your answer.

The overall rate law will match the rate law of the slowest step. The rate law of an elementary step can be determined by the reactants present, and in this case, the rate law for Step 1 matches the overall rate law. Therefore, Step 1 is the slowest step.

(ii) If the reaction is measured over the course of several minutes, what would the units of the rate constant be?

Using unit analysis:

$$\text{rate} = k[\text{NH}_3][\text{OCl}^-]$$

$$M/\text{min} = k(M)(M)$$

$$k = M^{-1}\text{min}^{-1}$$

$$8. (a) (i) \frac{(3.8 \times 10^{-5} M) - (2.6 \times 10^{-5} M)}{60 \text{ s} - 120 \text{ s}} = 2.0 \times 10^{-7} M/\text{s}$$

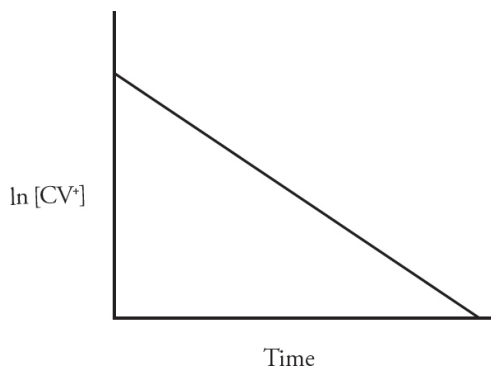
(ii) The solution starts reacting (and thus, fading) immediately after it is mixed. If the student waited 30 seconds before putting the cuvette in, the calculated rate of disappearance would thus decrease.

(b) $A = abc$

$$A = (26,000 \text{ cm}^{-1}\text{M}^{-1})(1.00 \text{ cm})(3.8 \times 10^{-5} M)$$

$$A = 0.98$$

(c) A reaction that is first order with respect to $[\text{CV}^+]$ will create a straight line in a graph of $\ln [\text{CV}^+]$ vs. time.



- (d) Between trials 1 and 2, the value of the $[CV^+]$ remained constant, while the $[OH^-]$ doubled. At the same time, the rate of reaction also doubled, meaning there is a direct relationship between $[OH^-]$ and the rate, so the reaction is first order with respect to $[OH^-]$. The rate with respect to $[CV^+]$ has already been established as first order, so:

$$\text{rate} = k[CV^+][OH^-]$$

- (e) Either of the trials can be used for this. Taking trial 2:

$$7.20 \times 10^{-7} \text{ M/s} = k(5.5 \times 10^{-5} \text{ M})(0.24 \text{ M})$$

$$k = 0.055 \text{ M}^{-1}\text{s}^{-1}$$

- (f) $\text{Rate} = k[CV^+][OH^-]$

$$\text{Rate} = (0.055 \text{ M}^{-1}\text{s}^{-1})(4.1 \times 10^{-5} \text{ M})(0.18 \text{ M})$$

$$\text{Rate} = 4.1 \times 10^{-7} \text{ s}^{-1}$$

Chapter 7

Big Idea #5: Laws of Thermodynamics and Changes in Matter

The laws of thermodynamics describe the essential role of energy and explain and predict the direction of changes in matter.

HEAT AND TEMPERATURE

Many people do not understand that heat and temperature are not, in fact, synonyms. The fact is, while the concepts are certainly related, they represent different things. **Temperature** represents the average amount of kinetic energy present in a given substance. **Heat** represents energy flow between two substances at different temperatures. Energy can be neither created nor destroyed; the amount of energy in the universe is constant. This is sometimes called the **First Law of Thermodynamics**.

Energy transfers due to molecular collisions. As the faster moving molecules collide with the slower moving ones, they transfer some of their energy, changing the speed of both molecules. We've already defined the terms exothermic and endothermic in terms of enthalpy back in chapter 5, but those terms can also be defined another way when studying the energy flow that takes place with many physical and chemical processes. In an exothermic process, energy is transferred from the reaction itself to its surroundings. If you dissolve salt in water and the temperature of the water increases, it is because the process itself (the salt dissolving in water) emitted energy. The surroundings (in this case, the water), absorbed it.

For an endothermic process the opposite is true. Energy transfers from the surroundings into the process. When an ice cube melts in your hand, the energy leaves your hand (and the surrounding air) and goes into the ice cube. This is an endothermic process. Both physical and chemical changes can be classified as exothermic or endothermic.

STATE FUNCTIONS

Enthalpy change (ΔH), entropy change (ΔS), and free-energy change (ΔG) are **state functions**. That means they all depend only on the change between the initial and final states of a system, not on the process by which the change occurs. For a chemical reaction, this

means that the thermodynamic state functions are independent of the reaction pathway; for instance, the addition of a catalyst to a reaction will have no effect on the overall energy or entropy change of the reaction.

Standard State Conditions

When the values of thermodynamic quantities are given on the test, they are almost always given for standard state conditions. A thermodynamic quantity under standard state conditions is indicated by the little superscript circle, so the following is true under standard state conditions:

$$\Delta H = \Delta H^\circ$$

$$\Delta S = \Delta S^\circ$$

$$\Delta G = \Delta G^\circ$$

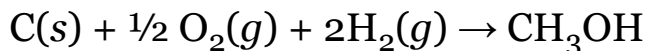
Standard State Conditions

- All gases are at 1 atmosphere pressure.
- All liquids are pure.
- All solids are pure.
- All solutions are at 1-molar (1 *M*) concentration.
- The energy of formation of an element in its normal state is defined as zero.
- The temperature used for standard state values is almost invariably room temperature: 25°C (298 K). Standard state values can be calculated for other temperatures, however.

ENTHALPY OF FORMATION, ΔH_f°

Enthalpy of formation (also known as heat of formation) is the change in energy that takes place when one mole of a compound is formed from its component pure elements under standard state conditions. Enthalpy of formation is almost always calculated at a temperature of 25°C (298 K).

An example of an enthalpy of formation equation is below.



Note that in any enthalpy of formation equation, there is always exactly one mole of product. When that product (in this case, CH₃OH, also known as methanol) contains an odd number of atoms from an element that is diatomic in nature, it is thus common practice to use halves to balance the equation. This practice of using fractions to balance equations is only used when dealing with diatomics and even then the only fractions that can be used are halves, so don't start getting creative when balancing other equations!

An easy mnemonic to remember the diatomic elements is Dr. HOBrINCl! (Hydrogen, oxygen, fluorine, bromine, iodine, nitrogen, and chlorine).

ΔH_f° for a pure element is defined as zero. This is even true for elements that, in their pure state, appear as diatomic molecules (such as oxygen and hydrogen).

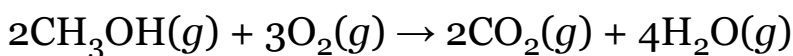
- If ΔH_f° for a compound is negative, energy is released when the compound is formed from pure elements, and the product is *more* stable than its constituent elements. That is, the process is exothermic.
- If ΔH_f° for a compound is positive, energy is absorbed when the compound is formed from pure elements, and the product is *less*

stable than its constituent elements. That is, the process is endothermic.

If the ΔH_f° values of the products and reactants are known, ΔH for a reaction can be calculated.

$$\Delta H^\circ = \Sigma \Delta H_f^\circ \text{ products} - \Sigma \Delta H_f^\circ \text{ reactants}$$

The ΔH for a reaction describes the energy change in that reaction when it goes to completion as written. On the AP Exam, the units for reaction enthalpy are kJ/mol_{rxn}. These units are often simply just written as kJ/mol, but either way they indicate the same quantity—the amount of heat released (negative ΔH) or absorbed (positive ΔH) when the reaction occurs with the mole quantities represented in the balanced equation. Let's find ΔH° for the reaction below.



Compound	(kJ/mol)
CH ₃ OH(<i>g</i>)	−201
O ₂ (<i>g</i>)	0
CO ₂ (<i>g</i>)	−394
H ₂ O(<i>g</i>)	−242

$$\Delta H^\circ = \Sigma \Delta H_f^\circ \text{ products} - \Sigma \Delta H_f^\circ \text{ reactants}$$

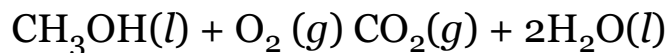
$$\Delta H^\circ = [(2)(\Delta H_f^\circ \text{ CO}_2) + (4)(\Delta H_f^\circ \text{ H}_2\text{O})] - [(2)(\Delta H_f^\circ \text{ CH}_3\text{OH}) + (3)(\Delta H_f^\circ \text{ O}_2)]$$

$$\Delta H^\circ = [(2)(-394 \text{ kJ}) + (4)(-242 \text{ kJ})] - [(2)(-201 \text{ kJ}) + (3)(0 \text{ kJ})]$$

$$\Delta H^\circ = (-1,756 \text{ kJ}) - (-402 \text{ kJ})$$

$$\Delta H^\circ = -1,354 \text{ kJ/mol}_{\text{rxn}}$$

In the example above, when 2 moles of CH_3OH react with 3 moles of oxygen, exactly 1354 kJ of energy is released. On a related note, if we were to look at HALF of the above overall reaction, we'd get:



$$\Delta H^\circ = -677 \text{ kJ/mol}$$

This represents the **enthalpy of combustion** for methanol. Combustion is always an exothermic process, and the enthalpy of combustion specifically describes the amount of energy released when ONE mole of a hydrocarbon combusts. Thus, when talking about enthalpy of combustion, we again often see halves used to balance the diatomic oxygen molecule that is always part of a combustion reaction. Note that the enthalpy change is exactly half of the calculated enthalpy change from above, as we are combusting exactly half as much methanol. However, when reactions occur, they are rarely in exact quantities like that. Instead, you have to combine stoichiometry concepts with thermodynamics to determine the energy change.

Example:

How much heat is released when 5.00 g of CH_3OH is combusted in excess oxygen?

In this case, the prompt tells us that CH_3OH is the limiting reactant. Otherwise, we would have to determine which reactant was limiting. The limiting reactant not only limits the amount of products formed, but it also limits the amount of heat change that occurs during the reaction.

Since we know the CH_3OH is limiting, we can set up some stoichiometry to get to our answer:

$$5.00 \text{ g CH}_3\text{OH} \times \frac{1 \text{ mol CH}_3\text{OH}}{32.05 \text{ g CH}_3\text{OH}} \times \frac{1 \text{ mol}_{\text{rxn}}}{2 \text{ mol CH}_3\text{OH}} \times \frac{-1354 \text{ kJ}}{1 \text{ mol}_{\text{rxn}}} = -106 \text{ kJ}$$

So, 106 kJ of energy is released (indicated by the negative sign). Don't forget to take mole ratios into account when doing thermodynamics problems that involve stoichiometry. As a general rule, whenever you are given specific amounts of reactants (either in grams or moles) and asked about the energy change for a reaction, you will have to use stoichiometry to get the correct answer.

BOND ENERGY

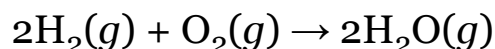
Bond energy is the energy required to break a bond. Because the breaking of a bond is an endothermic process, bond energy is always a positive number. When a bond is formed, energy equal to the bond energy is released.

$$\Delta H^\circ = \Sigma \text{ Bond energies of bonds broken} - \Sigma \text{ Bond energies of bonds formed}$$

The bonds broken will be the reactant bonds, and the bonds formed will be the product bonds.

The number of bonds broken and formed is affected by both the number of that particular type of bond within a molecule, as well as how many of those molecules there are in a balanced reaction. For instance, in the equation below, there are four O–H bonds being formed because each water molecule has two O–H bonds, and there are two water molecules present.

Let's find ΔH° for the following reaction.



Bond

Bond Energy (kJ/mol)

H-H	436
O=O	499
O-H	463

$\Delta H^\circ = \Sigma \text{ Bond energies of bonds broken} - \Sigma \text{ Bond energies of bonds formed}$

$$\Delta H^\circ = [(2)(\text{H-H}) + (1)(\text{O=O})] - [(4)(\text{O-H})]$$

$$\Delta H^\circ = [(2)(436 \text{ kJ}) + (1)(499 \text{ kJ})] - [(4)(463 \text{ kJ})]$$

$$\Delta H^\circ = (1,371 \text{ kJ}) - (1,852 \text{ kJ})$$

$$\Delta H^\circ = -481 \text{ kJ/mol}$$

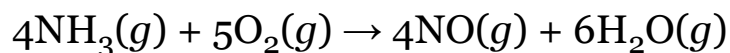
HESS'S LAW

Hess's Law states that if a reaction can be described as a series of steps, then ΔH for the overall reaction is simply the sum of the ΔH values for all the steps.

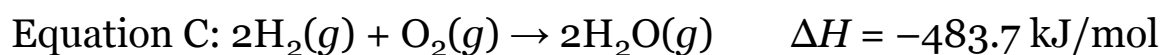
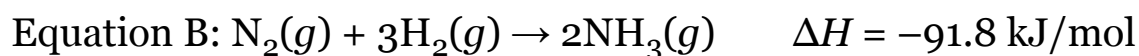
When manipulating equations for use in enthalpy calculations, there are three rules:

1. If you flip the equation, flip the sign on the enthalpy value.
2. If you multiply or divide an equation by an integer, also multiply/divide the enthalpy value by that same integer.
3. If several equations, when summed up, create a new equation, you can also add the enthalpy values of those component equations to get the enthalpy value for the new equation.

For example, let's say you want to calculate the enthalpy change for the following reaction:

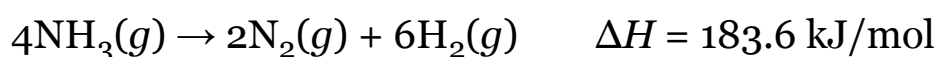
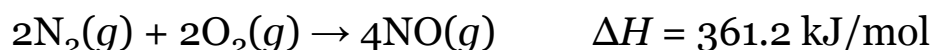


Given:



First, we want to make sure all of the equations have the products and reactants on the correct side. A quick glance shows us the NH_3 should end up on the left but is given to us on the right, so equation B needs to be flipped, which will change the ΔH value to 91.8 kJ/mol . All other species appear to be on the side they need to be on, so we'll leave the other two reactions alone.

Next, we're going to see what coefficients we need to get to. The NO needs to have a 4 but is only 2 in equation A, so we'll multiply that equation by 2, which will change the ΔH value to $180.6 \times 2 = 361.2 \text{ kJ/mol}$. The NH_3 also needs to have a 4 but is only 2 in equation B, so we'll also multiply that equation by 2, changing ΔH to $91.8 \times 2 = 183.6 \text{ kJ/mol}$. Finally, the H_2O needs to have a 6 but only has a 2 in equation C. We now have:



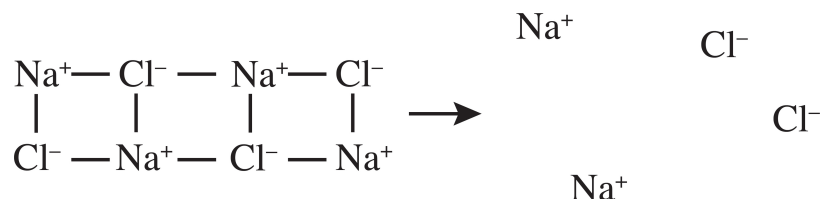
Fortunately, the two O_2 on the products side add up to 5, and both the N_2 and H_2 cancel out completely, giving us the final equation that we wanted. So, our final value for the enthalpy will be $361.2 + 183.6 - 1451 = -906 \text{ kJ/mol}$.

ENTHALPY OF SOLUTION

When an ionic substance dissolves in water, a certain amount of heat is emitted or absorbed. The bond between the cation and anion is breaking, which requires energy, but energy is released when those ions form new attractions to the dipoles of the water molecules.

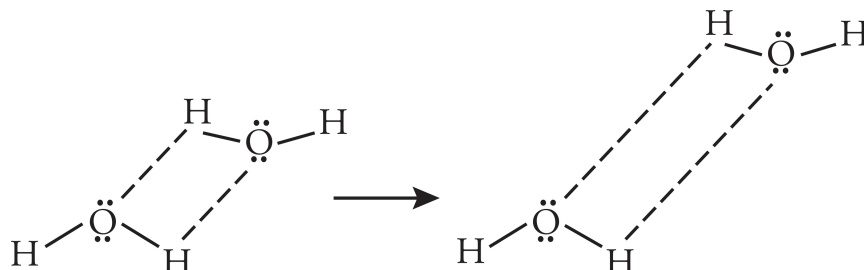
Typically, this process can be broken down into three steps. We will use NaCl dissolving in water for this example.

Step 1: Breaking the Solute Bonds



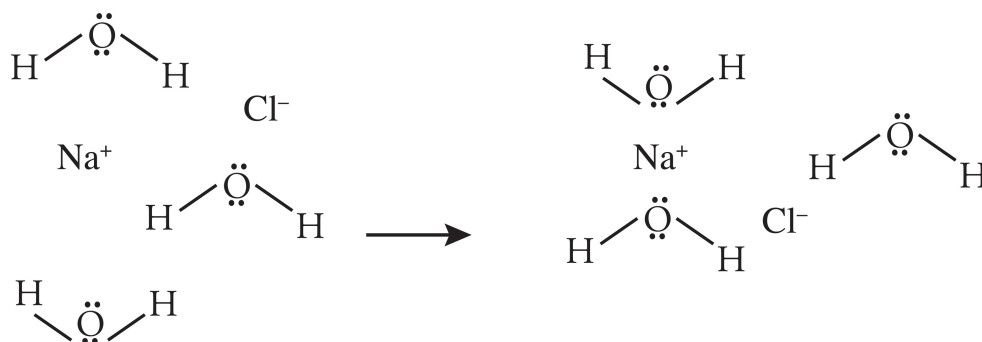
The bonds between the Na^+ and Cl^- ions must be broken. The amount of energy needed to break this bond is equal to the lattice energy (see [this page](#) for a more detailed explanation on lattice energy). As energy is required to break the bonds, this step always has a positive ΔH .

Step 2: Separating the Solvent Molecules



The water molecules must spread out to make room for the Na^+ and Cl^- ions. This requires energy to weaken (but not break) the intermolecular forces between them, so this step always has a positive ΔH .

Step 3: Creating New Attractions



The last step involves the free-floating ions being attracted to the dipoles of the water molecules. Note that while bonds are not being formed (no electrons are transferring or being shared), energy is still released in this process. As such, this step always has a negative ΔH .

The energy values from step 2 and step 3 combined are often called the **hydration energy**. This value is always negative, as the magnitude of the energy change in step 3 exceeds the magnitude of the energy change in step 2. As with lattice energy, hydration energy is a Coulombic energy and thus increases as the ions either increase in charge (i.e., $\text{Mg}^{2+} > \text{Na}^+$) or decrease in size (i.e., $\text{Na}^+ > \text{K}^+$).

If you add the energy values for all three steps together, you can determine the enthalpy of solution for that compound. Thus, if the hydration energy exceeds the lattice energy, the enthalpy of solution is negative. However, if the lattice energy exceeds the hydration energy, the enthalpy of solution is positive.

THERMODYNAMICS OF PHASE CHANGE

Naming the Phase Changes

Phase changes occur because of changes in temperature and/or pressure. Phase changes are always physical changes, as they do not involve breaking any bonds or the creation of new substances.

Some particles in a liquid or solid will have enough energy to break away from the surface and become gaseous. The pressure exerted by these molecules as they escape from the surface is called the **vapor**

pressure. When the liquid or solid phase of a substance is in equilibrium with the gas phase, the pressure of the gas will be equal to the vapor pressure of the substance. As temperature increases, the vapor pressure of a liquid will increase. When the vapor pressure of a liquid increases to the point where it is equal to the surrounding atmospheric pressure, the liquid boils.

<i>Solid to liquid</i>	— Melting
<i>Liquid to solid</i>	— Freezing
<i>Liquid to gas</i>	— Vaporization
<i>Gas to liquid</i>	— Condensation
<i>Solid to gas</i>	— Sublimation
<i>Gas to solid</i>	— Deposition

Heat of Fusion

The **heat of fusion** is the energy that must be put into a solid to melt it. This energy is needed to overcome the forces holding the solid together. Alternatively, the heat of fusion is the heat given off by a substance when it freezes. The intermolecular forces within a solid are more stable and therefore have lower energy than the forces within a liquid, so energy is released in the freezing process.

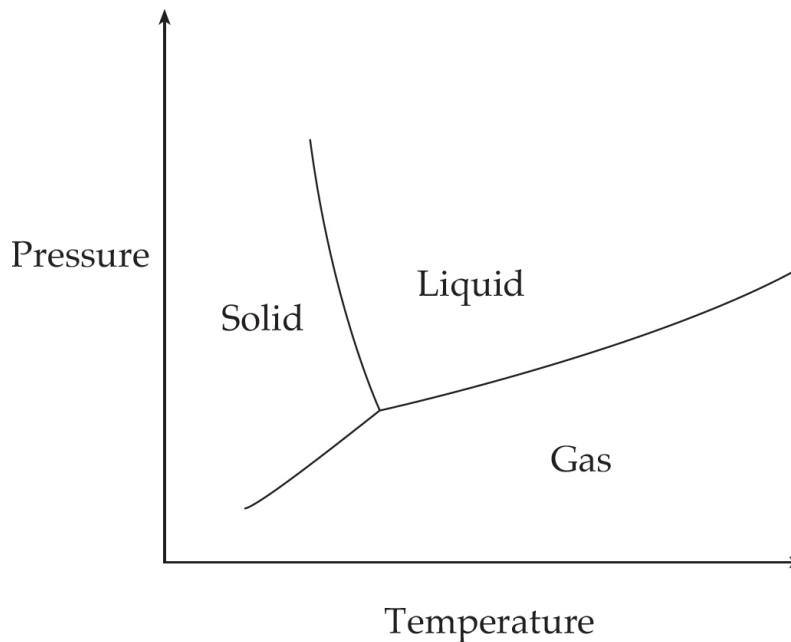
Heat of Vaporization

The **heat of vaporization** is the energy that must be put into a liquid to turn it into a gas. This energy is needed to overcome the forces holding the liquid together. Alternatively, the heat of vaporization is the heat given off by a substance when it condenses. Intermolecular forces become stronger when a gas condenses; the gas becomes a liquid, which is more stable, and energy is released.

As heat is added to a substance, the temperature of the substance can increase or the substance can change phases, but both changes cannot occur simultaneously. Therefore, when a substance is changing phases, the temperature of that substance remains constant.

Phase Diagram for Water

A phase diagram tells you the state of a substance at various pressure/temperature combinations. The phase diagram for water is below.



In the phase diagram for substances other than water, the solid-liquid equilibrium line slopes upward. In the phase diagram for water, the solid-liquid equilibrium line slopes *downward*. What this means is that when pressure is increased, a normal substance will change from liquid to solid, but water will change from solid to liquid.

Water has this odd property because its hydrogen bonds form a lattice structure when it freezes. This forces the molecules to remain farther apart in ice than in water, making the solid phase less dense

than the liquid phase. That's why ice floats on water, and this natural phenomenon preserves most marine life.

CALORIMETRY

The specific heat of a substance is the amount of heat required to raise the temperature of one gram (or kilogram) of that substance by one degree Celsius (or one Kelvin). An object with a large **specific heat** can absorb a lot of heat without undergoing much of a temperature change, while a substance with a low specific heat will experience much greater temperature changes.

The easiest way to think about this is by considering the temperature changes you experience when wearing a black shirt as opposed to a white shirt on a hot summer day. You are absorbing the same amount of heat from the sun regardless of which shirt you are wearing. However, the black shirt has a considerably lower specific heat than the white shirt, and thus will experience a greater temperature change—making you a lot hotter!

Specific Heat

$$q = mc\Delta T$$

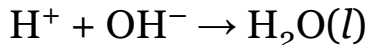
q = heat added (J or cal)

m = mass of the substance (g or kg)

c = specific heat

ΔT = temperature change (K or °C)

Calorimetry is the measurement of heat changes during chemical reactions, and it is frequently accomplished via the equation above. Determining the amount of heat transfer in a reaction can lead directly to determining the enthalpy for that reaction, as shown in the example below.



25.0 mL of 1.5 *M* HCl and 30.0 mL of 2.0 *M* NaOH are mixed together in a Styrofoam cup and the reaction above occurs. The temperature of the reaction rises from 23.00°C to 31.60°C over the course of the reaction. Assuming the density of the solutions is 1.0 g/mL and the specific heat of the mixture is 4.18 J/g°C, calculate the enthalpy of the reaction.

To solve this, we first need to determine the amount of heat released during the reaction. The mass of the final solution can be determined by taking the total volume of the solution (55.0 mL) and multiplying that by the density, yielding 55.0g. The temperature change is 31.60 – 23.00 = 8.60°C. So:

$$q = mc\Delta T$$

$$q = (55.0\text{g})(4.18 \text{ J/g}^\circ\text{C})(8.60^\circ\text{C})$$

$$q = 1970 \text{ J}$$

From here, we need to determine how many moles of product are formed in this reaction. Looking at the two reactants, we can see that there is going to be less of the HCl, which has a lower molarity and volume. As everything here is in a 1:1 ratio, that means the HCl will be limiting and we can use it to determine how many moles of product will form.

$$\text{Molarity} = \text{mol/volume}$$

$$1.5 \text{ M} = n/0.025 \text{ L}$$

$$n = 0.038 \text{ mol HCl} = 0.038 \text{ mol H}_2\text{O formed}$$

The calculated heat gain from earlier (1970 J) was the heat gained by the water. Due to conservation of energy, this is also the heat lost by the reaction itself. To calculate the enthalpy of reaction, we have to

flip that sign to indicate heat is lost, and then divide that value by the number of moles.

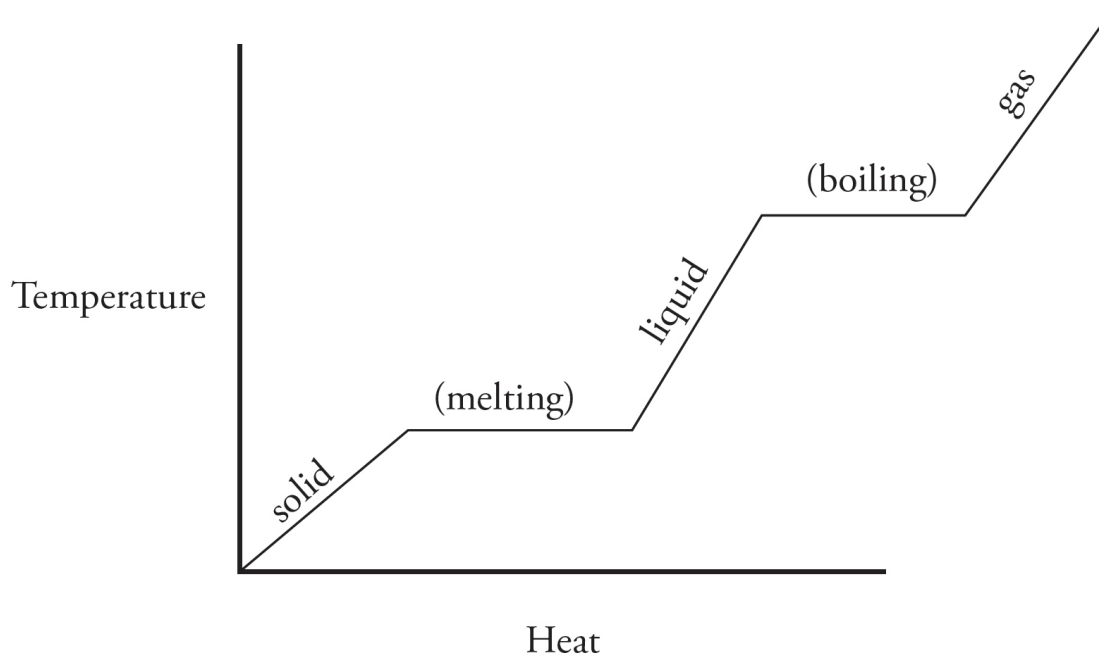
$$\Delta H = -1970 \text{ J}/0.038 \text{ mol} = -52,000 \text{ J/mol} = -52 \text{ kJ/mol}$$

It is important to emphasize that ΔH is always measured in joules per mole (or kilojoules per mole). Enthalpy is not just heat; it's the amount of heat given per mole of product that is created. Limiting reagent calculations can be done in situations where it is not easy to determine which reactant is limiting by inspection, as we did above.

Also, bear in mind that if you completely dissolve a solid into water, you must add the mass of that solid to the mass of the water when doing your calculations. So, if you were to dissolve 5.0 g of NaCl into 200.0 mL of water (remember, the density of water is 1.0 g/mL), the final mass of the solution would be 205.0 g.

HEATING CURVES

A heating (or cooling) curve shows what happens to the temperature of a substance as heat is added. If the substance is in a single phase, the temperature of the substance will increase. The amount of the temperature increase can be calculated using calorimetry. If the substance is in the process of a phase change, the temperature will remain constant and the amount of heat required to fully cause the substance to change phases can be calculated using the heat of fusion or the heat of vaporization.



There are a lot of different units that can be used when talking about these heat versus temperature curves, so it is very important to pay extremely close attention to what your units are when doing calculations.

Example:

A 1.53 g piece of ice is in a freezer and initially at a temperature of -15.1°C . The ice is removed from the freezer and melts completely after reaching a temperature of 0.0°C . If the specific heat of ice is $2.03 \text{ J/g}^{\circ}\text{C}$ and its molar heat of fusion is 6.01 kJ/mol , how much heat is required for the entire process to occur?

There are two parts to this problem. The first part involves the temperature change of the ice.

$$q = mc\Delta T$$

$$q = (1.53\text{g})(2.03 \text{ J/g}^{\circ}\text{C})(15.1^{\circ}\text{C})$$

$$q = 46.9 \text{ J}$$

The second part involves the ice melting. As this is a molar heat capacity, the number of moles of ice must first be determined.

$$1.53 \text{ g H}_2 \times \frac{1 \text{ mol H}_2\text{O}}{18.01 \text{ g}} = 0.0850 \text{ mol H}_2\text{O}$$

$$6.01 \text{ kJ/mol} \times 0.0850 \text{ mol} = 0.511 \text{ kJ} = 511 \text{ J}$$

Finally, the two values can be added together.

$$46.9 \text{ J} + 511 \text{ J} = 558 \text{ J}$$

ENTROPY

The entropy, S , of a system is a measure of the randomness or disorder of the system; the greater the disorder of a system, the greater its entropy. Because zero entropy is defined as a solid crystal at 0 K, and because 0 K has never been reached experimentally, all substances that we encounter will have some positive value for entropy. Standard entropies, S° , are calculated at 25°C (298 K).

The standard entropy change, ΔS° , that has taken place at the completion of a reaction is the difference between the standard entropies of the products and the standard entropies of the reactants.

$$\Delta S^\circ = \Sigma S^\circ_{\text{products}} - \Sigma S^\circ_{\text{reactants}}$$

You should be familiar with several simple rules concerning entropies.

- Liquids have higher entropy values than solids.
- Gases have higher entropy values than liquids.
- Particles in solution have higher entropy values than solids.

- Two moles of a gaseous substance have a higher entropy value than one mole of a gaseous substance.

GIBBS FREE ENERGY

The **Gibbs free energy**, or simply free energy, G , of a process is a measure of whether or not the process will proceed without the input of outside energy. A process that occurs without outside energy input is said to be thermodynamically favored, while one that does not is said to be thermodynamically unfavored. Prior to the 2014 exam, the terms *spontaneous* and *nonspontaneous* were used to describe thermodynamically favored and unfavored reactions, respectively. It would be good for you to be familiar with both sets of terms, although this text will use the updated terms.

FREE ENERGY CHANGE, ΔG

The standard free energy change, ΔG , for a reaction can be calculated from the standard free energies of formation, G°_f , of its products and reactants in the same way that ΔS° was calculated.

$$\Delta G^\circ = \Sigma \Delta G^\circ_{f \text{ products}} - \Sigma \Delta G^\circ_{f \text{ reactants}}$$

For a given reaction:

- if ΔG is negative, the reaction is thermodynamically favored
- if ΔG is positive, the reaction is thermodynamically unfavored
- if $\Delta G = 0$, the reaction is at equilibrium

ΔG , ΔH , AND ΔS

In general, nature likes to move toward two different and seemingly contradictory states—low energy and high disorder, so

thermodynamically favored processes must result in decreasing enthalpy or increasing entropy or both.

Nothing To See Here

Note that for all phase changes, no chemical reaction is occurring; therefore, the ΔG for any phase change is 0.

There is an important equation that relates favorability (ΔG), enthalpy (ΔH), and entropy (ΔS) to one another.

$$\Delta G^\circ = \Delta H^\circ - T\Delta S^\circ$$

T = absolute temperature (K)

Note that you should make sure your units always match up here. Frequently, ΔS is given in J/mol•K and ΔH is given in kJ/mol. The convention is to convert the $T\Delta S$ term to kilojoules (kJ), as that is what Gibbs free energy is usually measured in. However, you can also convert the ΔH term to joules instead, so long as you are keeping track and labeling all units appropriately.

The chart below shows how different values of enthalpy and entropy affect spontaneity.

ΔH	ΔS	T	ΔG	
–	+	Low	–	Always favored
		High	–	
+	–	Low	+	Never favored
		High	+	
+	+	Low	+	Not favored at low

				temperature
		High	–	Favored at high temperature
–	–	Low	–	Favored at low temperature
		High	+	Not favored at high temperature

You should note that at low temperature, enthalpy is dominant, while at high temperature, entropy is dominant.

VOLTAGE AND FAVORABILITY

A redox reaction will be favored if its potential has a positive value. We also know from thermodynamics that a reaction that is favored has a negative value for free-energy change. The relationship between reaction potential and free energy for a redox reaction is given by the equation below, which serves as a bridge between thermodynamics and electrochemistry.

$$\Delta G^{\circ} = -nFE^{\circ}$$

ΔG° = Standard Gibbs free energy change
(J/mol)

n = the number of moles of electrons
exchanged in the reaction (mol)

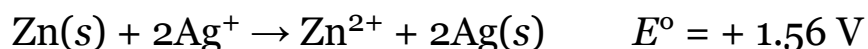
F = Faraday's constant, 96,500
coulombs/mole (that is, 1 mole of
electrons has a charge of 96,500
coulombs)

E° = Standard reaction potential (V, which
is equivalent to J/C)

From this equation we can see a few important things. If E° is positive, ΔG° is negative and the reaction is thermodynamically favored, and if E° is negative, ΔG° is positive and the reaction is thermodynamically unfavored.

Let's take a look at an example.

Calculate the ΔG value for the below reaction under standard conditions:



For this reaction, two moles of electrons are being transferred as the silver is reduced and the zinc is oxidized. So, $n = 2$. With that in mind:

$$\Delta G^\circ = - (2)(96500)(1.56)$$

$$\Delta G^\circ = - 301,000 \text{ J/mol}$$

As the ΔG° value is negative, we would expect this reaction to be favored under standard conditions, which is in line with the positive value for the cell.

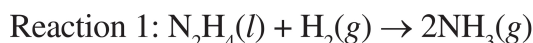
Note that your units on the free energy are in J/mol and not kJ/mol; this will always be the case with this equation, as one volt is equivalent to one joule/coulomb, so the answer comes out in joules instead of kilojoules.

CHAPTER 7 QUESTIONS

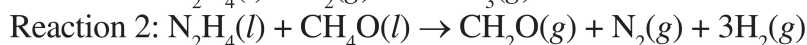
[Click here](#) to download and print the PDF version of this exercise.

Multiple-Choice Questions

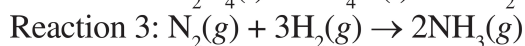
Use the following information to answer questions 1-5.



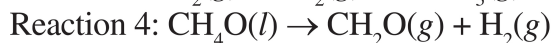
$$\Delta H = ?$$



$$\Delta H = -37 \text{ kJ/mol}_{\text{rxn}}$$



$$\Delta H = -46 \text{ kJ/mol}_{\text{rxn}}$$



$$\Delta H = -65 \text{ kJ/mol}_{\text{rxn}}$$

1. What is the enthalpy change for reaction 1?

(A) $-148 \text{ kJ/mol}_{\text{rxn}}$

(B) $-56 \text{ kJ/mol}_{\text{rxn}}$

(C) $-18 \text{ kJ/mol}_{\text{rxn}}$

(D) $+148 \text{ kJ/mol}_{\text{rxn}}$

2. If reaction 2 were repeated at a higher temperature, how would the reaction's value for ΔG be affected?

(A) It would become more negative because entropy is a driving force behind this reaction.

(B) It would become more positive because the reactant molecules would collide more often.

(C) It would become more negative because the gases will be at a higher pressure.

(D) It will stay the same; temperature does not affect the value for ΔG .

3. Under what conditions would reaction 3 be thermodynamically favored?

- (A) It is always favored.
- (B) It is never favored.
- (C) It is only favored at low temperatures.
- (D) It is only favored at high temperatures.

4. If 64 g of CH_4O were to decompose via reaction 4, approximately how much energy would be released or absorbed?

- (A) 65 kJ of energy will be absorbed.
- (B) 65 kJ of energy will be released.
- (C) 130 kJ of energy will be absorbed.
- (D) 130 kJ of energy will be released.

5. A 2.0 L flask holds 0.40 g of helium gas. If the helium is evacuated into a larger container while the temperature is held constant, what will the effect on the entropy of the helium be?

- (A) It will remain constant because the number of helium molecules does not change.
- (B) It will decrease because the gas will be more ordered in the larger flask.
- (C) It will decrease because the molecules will collide with the sides of the larger flask less often than they did in the smaller flask.
- (D) It will increase because the gas molecules will be more dispersed in the larger flask.

6. $2\text{Al}(s) + 3\text{Cl}_2(g) \rightarrow 2\text{AlCl}_3(s)$

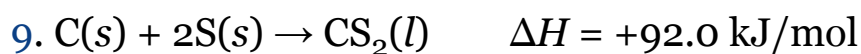
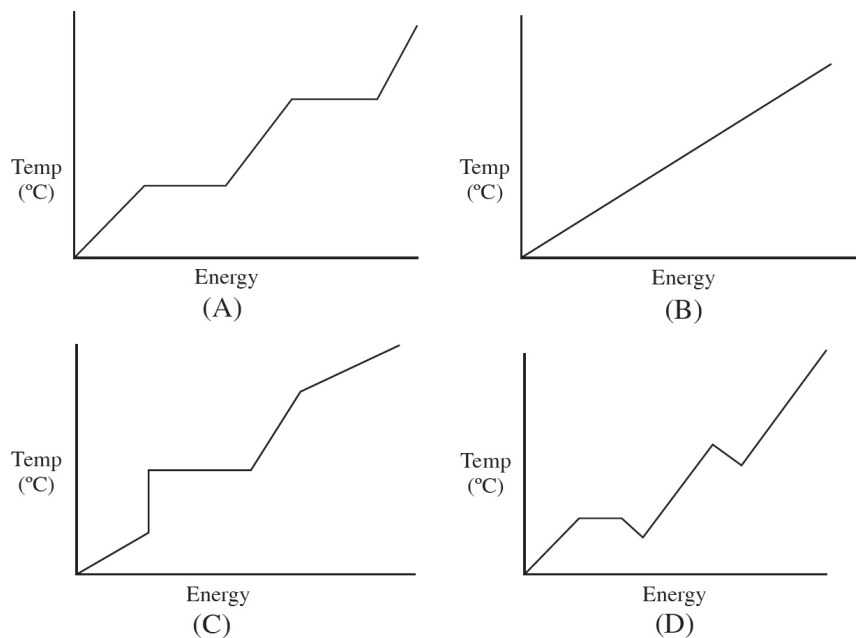
The reaction above is not thermodynamically favored under standard conditions, but it becomes thermodynamically favored as the temperature decreases toward absolute zero. Which of the following is true at standard conditions?

- (A) ΔS and ΔH are both negative.
- (B) ΔS and ΔH are both positive.
- (C) ΔS is negative, and ΔH is positive.
- (D) ΔS is positive, and ΔH is negative.

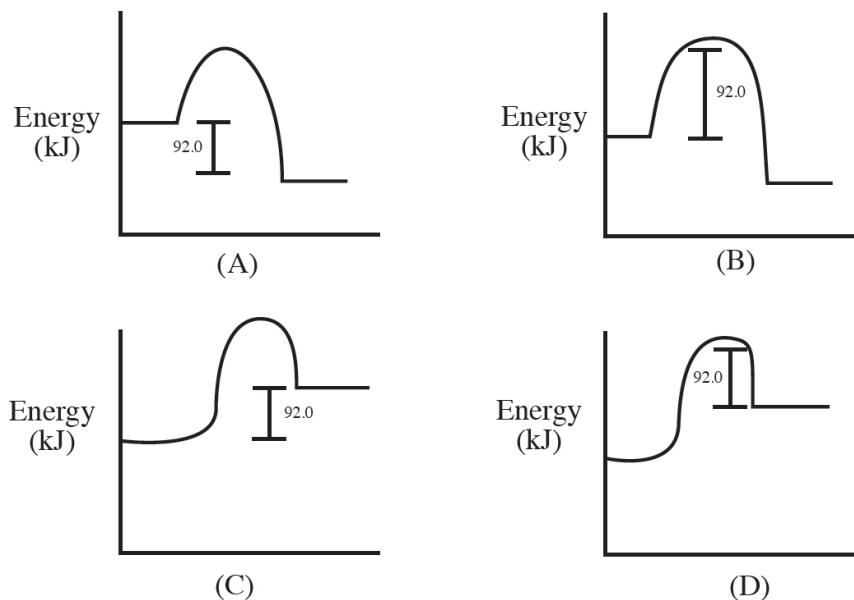
7. 1.50 g of NaNO_3 is dissolved into 25.0 mL of water, causing the temperature to increase by 2.2°C . The density of the final solution is found to be 1.02 g/mL. Which of the following expressions will correctly calculate the heat gained by the water as the NaNO_3 dissolves? Assume the volume of the solution remains unchanged.

- (A) $(25.0)(4.18)(2.2)$
- (B) $\frac{(26.5)(4.18)(2.2)}{1.02}$
- (C) $\frac{(1.02)(4.18)(2.2)}{1.50}$
- (D) $(25.0)(1.02)(4.18)(2.2)$

8. A pure solid substance is heated strongly. It first melts into a liquid, then boils and becomes a gas. Which of the following heating curves correctly shows the relationship between temperature and heat added?



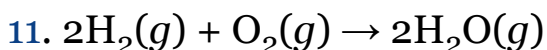
Which of the following energy level diagrams gives an accurate representation of the above reaction?



10. A student studies two solutions. Solution A has a volume of 100 mL and is at a temperature of 25.0°C. Solution B has a volume

of 1000 mL and is at a temperature of 22.0°C. Which of the following statements must be true regarding both solutions?

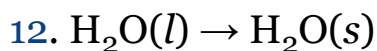
- (A) Solution A has more heat than solution B.
- (B) The specific heat capacity of solution A is greater than that of solution B.
- (C) If the solutions were to be mixed, heat would transfer from B to A.
- (D) Solution B has more thermal energy than solution A.



Based on the information given in the table below, what is ΔH° for the above reaction?

<u>Bond</u>	<u>Average bond energy (kJ/mol)</u>
H–H	500
O=O	500
O–H	500

- (A) –2,000 kJ
- (B) –500 kJ
- (C) +1,000 kJ
- (D) +2,000 kJ



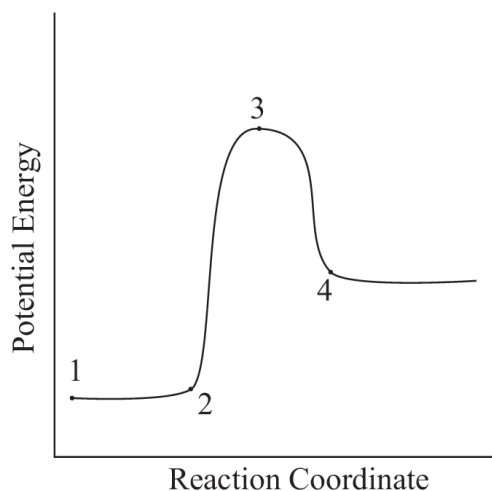
Which of the following is true for the above reaction?

- (A) The value for ΔS is positive.
- (B) The value for ΔG is zero.

(C) The value for ΔH is positive.

(D) The reaction is favored at 1.0 atm and 298 K.

Use the following diagram to answer questions 13-15.



13. Which point on the graph shown above corresponds to activated complex or transition state?

(A) 1

(B) 2

(C) 3

(D) 4

14. The distance between which two points is equal to the enthalpy change for this reaction?

(A) Points 1 and 2

(B) Points 1 and 3

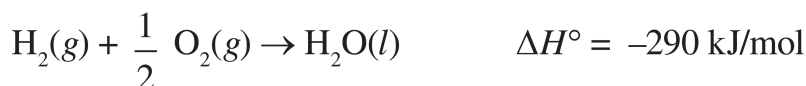
(C) Points 1 and 4

(D) Points 2 and 3

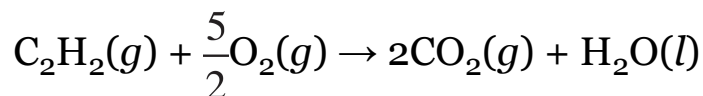
15. What would happen to this graph if a catalyst were to be added?

- (A) Point 3 would be lower.
- (B) The distance between points 2 and 4 would be decreased.
- (C) The slope of the line between points 2 and 3 would increase.
- (D) Point 4 would be higher.

16.



Based on the information given above, what is ΔH° for the following reaction?

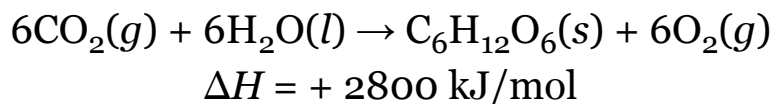


- (A) -1,300 kJ
- (B) -1,070 kJ
- (C) -840 kJ
- (D) -780 kJ

17. In which of the following reactions is entropy increasing?

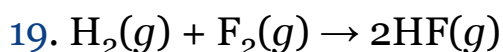
- (A) $2\text{SO}_2(g) + \text{O}_2(g) \rightarrow 2\text{SO}_3(g)$
- (B) $\text{CO}(g) + \text{H}_2\text{O}(g) \rightarrow \text{H}_2(g) + \text{CO}_2(g)$
- (C) $\text{H}_2(g) + \text{Cl}_2(g) \rightarrow 2\text{HCl}(g)$
- (D) $2\text{NO}_2(g) \rightarrow 2\text{NO}(g) + \text{O}_2(g)$

18. Consider the following reaction showing photosynthesis:



Which of the following is true regarding the thermal energy in this system?

- (A) It is transferred from the surroundings to the reaction.
- (B) It is transferred from the reaction to the surroundings.
- (C) It is transferred from the reactants to the products.
- (D) It is transferred from the products to the reactants.



Gaseous hydrogen and fluorine combine in the reaction above to form hydrogen fluoride with an enthalpy change of -540 kJ . What is the value of the heat of formation of $\text{HF}(g)$?

- (A) $-1,080 \text{ kJ/mol}$
- (B) -270 kJ/mol
- (C) 270 kJ/mol
- (D) 540 kJ/mol

Use the following information to answer questions 20-23.

When calcium chloride (CaCl_2) dissolves in water, the temperature of the water increases dramatically.

20. Which of the following must be true regarding the enthalpy of solution?

- (A) The lattice energy in CaCl_2 exceeds the bond energy within the water molecules.

- (B) The hydration energy between the water molecules and the solute ions exceeds the lattice energy within CaCl_2 .
- (C) The strength of the intermolecular forces between the solute ions and the dipoles on the water molecules must exceed the hydration energy.
- (D) The hydration energy must exceed the strength of the intermolecular forces between the water molecules.

21. During this reaction, heat transfers from

- (A) the reactants to the products
- (B) the reactants to the system
- (C) the system to the surroundings
- (D) the products to the surroundings

22. Which is the primary driving factor behind this reaction?

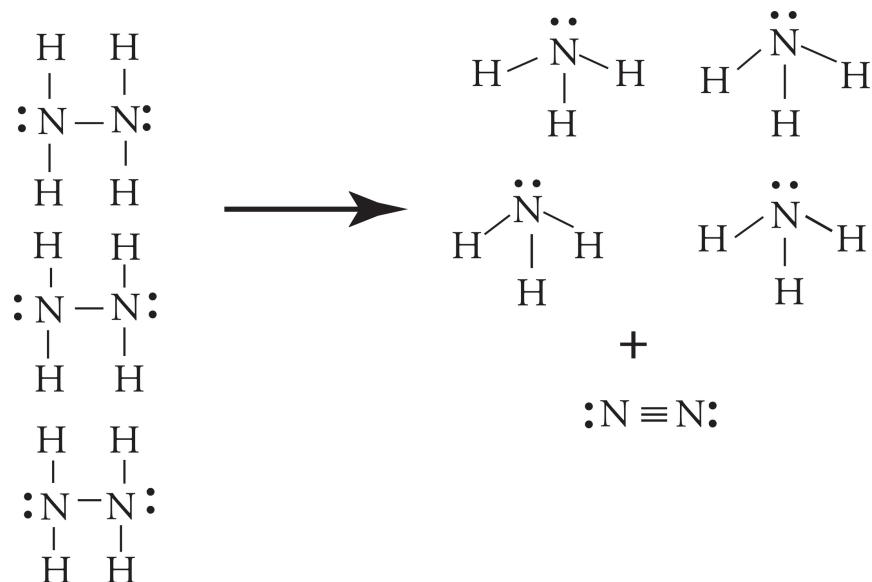
- (A) Entropy
- (B) Enthalpy
- (C) Both enthalpy and entropy
- (D) Neither enthalpy nor entropy

23. Compared to CaCl_2 , what must be true regarding the hydration energy of CaF_2 ?

- (A) It would be greater because fluoride is smaller than chloride.
- (B) It would be the same because the charges of fluoride and chloride are identical.
- (C) It would be the same because hydration energy is only dependent on the IMFs present in water.

(D) It would be smaller because the molar mass of CaF_2 is smaller than that of CaCl_2 .

24. The reaction shown in the diagram below is accompanied by a large increase in temperature. If all molecules shown are in their gaseous state, which statement accurately describes the reaction?



- (A) It is an exothermic reaction in which entropy increases.
(B) It is an exothermic reaction in which entropy decreases.
(C) It is an endothermic reaction in which entropy increases.
(D) It is an endothermic reaction in which entropy decreases.

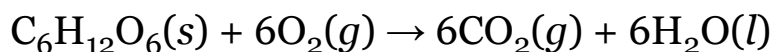
Free-Response Questions

1.

Substance	Absolute Entropy, S° (J/mol•K)	Molar Mass (g/mol)
$\text{C}_6\text{H}_{12}\text{O}_6(s)$	212.13	180

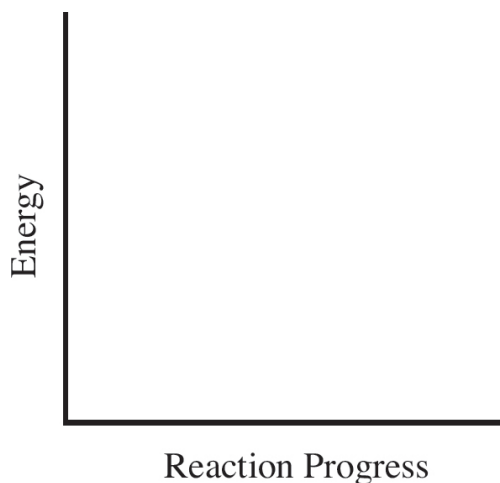
O ₂ (g)	205	32
CO ₂ (g)	213.6	44
H ₂ O(l)	69.9	18

Energy is released when glucose is oxidized in the following reaction, which is a metabolism reaction that takes place in the body.



The standard enthalpy change, ΔH° , for the reaction is $-2,801 \text{ kJ/mol}_{\text{rxn}}$ at 298 K.

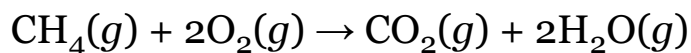
- Calculate the standard entropy change, ΔS° , for the oxidation of glucose.
- Calculate the standard free energy change, ΔG° , for the reaction at 298 K.
- Using the axis below, draw an energy profile for the reaction and indicate the magnitude of ΔH .



- How much energy is given off by the oxidation of 1.00 gram of glucose?

2.

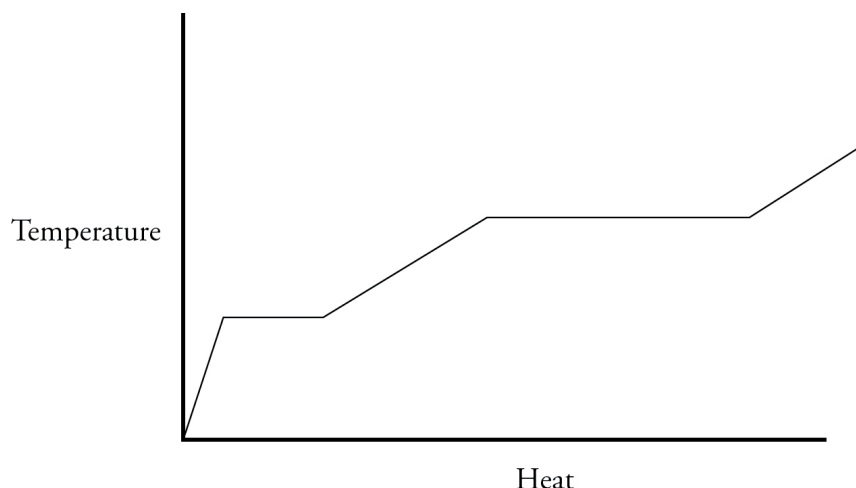
Bond	Average Bond Dissociation Energy (kJ/mol)
C–H	415
O=O	495
C=O	799
O–H	463



The standard free energy change, ΔG° , for the reaction above is $-801 \text{ kJ/mol}_{\text{rxn}}$ at 298 K.

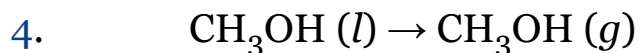
- (a) Use the table of bond dissociation energies to find ΔH° for the reaction above.
- (b) How many grams of methane must react with excess oxygen in order to release 1500 kJ of heat?
- (c) What is the value of ΔS° for the reaction at 298 K?
- (d) Give an explanation for the size of the entropy change found in (c).

3.



The diagram above shows how the temperature of a certain covalent substance changes as heat is added to it.

- (a) Which is greater for the substance: the heat of fusion or the heat of vaporization? How do you know?
- (b) If additional heat is added to the substance, the line would continue at its current slope and never become horizontal again. Why is this?
- (c) Reading the graph above, a student theorizes that the specific heat capacity of the substance is greatest in the solid phase. Do you agree? Why or why not?
- (d) What would the signs be for the enthalpy and entropy changes as heat continues to be added? Justify your answer.

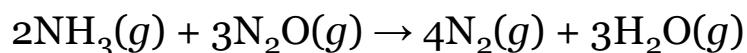


For the boiling of methanol, CH_3OH , $\Delta H = +37.6 \text{ kJ/mol}$ and $\Delta S = +111 \text{ J/mol} \cdot \text{K}$.

- (a)
 - (i) Why is the ΔH value positive for this process?
 - (ii) Why is the ΔS value positive for this process?
- (b) What is the boiling point of methanol in degrees Celsius?

- (c) How much heat is required to boil 50.0 mL of ethanol if the density of ethanol is 0.789 g/mL?
- (d) What will happen to the temperature of the methanol as it boils? Explain.
- (e) Would methanol be soluble with water? Why or why not?
- (f) Would you expect the boiling point of ethanol, $\text{CH}_3\text{CH}_2\text{OH}$, to be less than, greater than, or the same as methanol? Justify your answer.

5. Ammonia gas reacts with dinitrogen monoxide via the following reaction:



The absolute entropy values for the varying substances are listed in the table below.

Substance	S° (J/mol·K)
$\text{NH}_3(g)$	193
$\text{N}_2\text{O}(g)$	220
$\text{N}_2(g)$	192
$\text{H}_2\text{O}(g)$	189

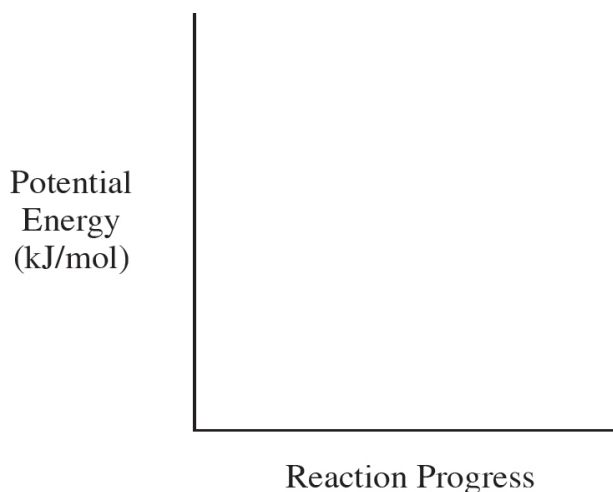
- (a) Calculate the entropy value for the overall reaction.

Several bond enthalpies are listed in the table below.

Bond	Enthalpy (kJ/mol)	Bond	Enthalpy (kJ/mol)
N–H	388	N=N	409
N–O	210	N≡N	941

N=O	630	O-H	463
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- (b) Calculate the enthalpy value for the overall reaction.
- (c) Is this reaction thermodynamically favored at 25°C? Justify your answer.
- (d) If 25.00 g of NH₃ reacts with 25.00 g of N₂O:
- Will energy be released or absorbed?
 - What is the magnitude of the energy change?
- (e) On the reaction coordinates below, draw a line showing the progression of this reaction. Label both ΔH and E_a on the graph.



6. A student designs an experiment to determine the specific heat of aluminum. The student heats a piece of aluminum with a mass of 5.86 g to various temperatures, then drops it into a calorimeter containing 25.0 mL of water. The following data is gathered during one of the trials:

Initial Temperature of Al (°C)	Initial Temperature of H ₂ O (°C)	Final Temperature of Al + H ₂ O (°C)

109.1	23.2	26.8
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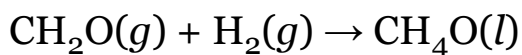
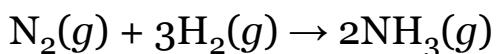
- (a) Given that the specific heat of water is $4.18 \text{ J/g} \times ^\circ\text{C}$ and assuming its density is exactly 1.00 g/mL , calculate the heat gained by the water.
- (b) Calculate the specific heat of aluminum from the experimental data given.
- (c) Calculate the enthalpy change for the cooling of aluminum in water in kJ/mol .
- (d) If the accepted specific heat of aluminum is $0.900 \text{ J/g} \times ^\circ\text{C}$, calculate the percent error.
- (e) Suggest two potential sources of error that would lead the student's experimental value to be different from the actual value. Be specific in your reasoning and make sure any identified error can be quantitatively tied to the student's results.

CHAPTER 7 ANSWERS AND EXPLANATIONS

Multiple-Choice

1. C

To get reaction 1, all that is needed is to flip reaction 4 and then add it to reactions 2 and 3.



$$(-37 \text{ kJ/mol}_{\text{rxn}}) + (-46 \text{ kJ/mol}_{\text{rxn}}) + (65 \text{ kJ/mol}_{\text{rxn}}) = -18 \text{ kJ/mol}_{\text{rxn}}$$

2. A

Entropy is positive during reaction 2, as creating gas molecules out of liquid molecules demonstrates an increase in disorder. Using $\Delta G = \Delta H - T\Delta S$, if the temperature increases, the $T\Delta S$ term will also increase. As the overall value for that term is negative, increasing $T\Delta S$ makes ΔG more negative.

3. C

Use the $\Delta G = \Delta H - T\Delta S$ equation. To do that, determine the signs of both ΔH and the $T\Delta S$ term.

You know that ΔH is negative. ΔS is negative as well, because the reaction is going from four molecules to two molecules, meaning it is becoming more ordered. That, in turn, means the $T\Delta S$ term is positive.

Reactions are only favored when ΔG is negative. As temperature increases, the $T\Delta S$ term becomes larger and more likely to be greater in magnitude than the ΔH term. If the temperature remains low, the $T\Delta S$ value is much more likely to be smaller in magnitude than the ΔH value, meaning ΔG is more likely to be negative and the reaction will be favored.

4. **D**

The molar mass of CH_4O is 32 g/mol, so 64 g of it represents two moles. For every one mole of CH_4O that decomposes, 65 kJ of energy is released—this is indicated by the negative sign on the H value. If two moles decompose, twice that amount—130 kJ—will be released.

5. **D**

Entropy is a measure of a system's disorder. In a larger flask, the gas molecules will spread farther apart and become more disordered.

6. **A**

Remember that $\Delta G = \Delta H - T\Delta S$.

If the reaction is thermodynamically favored only when the temperature is very low, then ΔG is negative only when T is very small. This can happen only when ΔH is negative and ΔS is negative. A very small value for T will eliminate the influence of ΔS .

7. **D**

Use the formula $q = mc\Delta T$. The mass is the solution's mass, which can be obtained by multiplying 25.0 mL by 1.02 g/mL.

8. **A**

During a phase change, all of the energy added goes toward breaking the intermolecular forces holding the molecules together. During this time, the speed of the molecules (and thus the temperature) does not rise.

9. **C**

A positive H means the reaction is endothermic, so the products have more bond energy than the reactants. The difference between the energy levels of the products and reactants is equal to ΔH . (The difference between the energy level of the reactants and the top of the hump is the value for the activation energy.)

10. **D**

Heat is a measure of energy transfer, so a solution cannot “have” heat. Solutions can have thermal energy, though, and even though solution B is at a lower temperature, there is a lot more of it, and the amount of energy contained within it will thus be greater.

11. **B**

The bond energy is the energy that must be put into a bond to break it. First, figure out how much energy must be put into the reactants to break their bonds.

To break 2 moles of H–H bonds, it takes $(2)(500) \text{ kJ} = 1,000 \text{ kJ}$.

To break 1 mole of O=O bonds, it takes 500 kJ.

So to break up the reactants, it takes +1,500 kJ.

Energy is given off when a bond is formed; that's the negative of the bond energy. Now see how much energy is given off when 2 moles of H_2O are formed.

2 moles of H_2O molecules contain 4 moles of O–H bonds, so $(4)(-500) \text{ kJ} = -2,000 \text{ kJ}$ are given off.

So the value of ΔH° for the reaction is

$(-2,000 \text{ kJ, the energy given off}) + (1,500 \text{ kJ, the energy put in}) = -500 \text{ kJ}$.

12. **B**

During a phase change, the products and reactants are at equilibrium with each other, meaning ΔG is zero. When a liquid turns into a solid, disorder decreases (negative ΔS) and energy is released (negative ΔH). Choice (D) is incorrect because $298 \text{ K} = 25^\circ\text{C}$, and water does not freeze at that temperature at 1.0 atm.

13. **C**

Point 3 represents the activated complex, which is the point of highest energy. This point is the transition state between the reactants and the products.

14. **C**

The enthalpy change is the distance between the energy level of the reactants (on which points 1 and 2 lie) and the energy level of the products (on which point 4 lies).

15. **A**

A catalyst lowers the activation energy of the reaction without changing the energy level of either the reactants or

products.

16. **A**

The equations given on top give the heats of formation of all the reactants and products (remember: the heat of formation of O_2 , an element in its most stable form, is zero).

ΔH° for a reaction = (ΔH° for the products) – (ΔH° for the reactants).

First, the products:

From CO_2 , you get $(2)(-390 \text{ kJ}) = -780 \text{ kJ}$.

From H_2O , you get -290 kJ .

So ΔH° for the products = $(-780 \text{ kJ}) + (-290 \text{ kJ}) = -1,070 \text{ kJ}$.

Now the reactants:

From C_2H_2 , you get $+230 \text{ kJ}$. The heat of formation of O_2 is defined as zero, so that's it for the reactants.

ΔH° for the reaction = $(-1,070 \text{ kJ}) - (+230 \text{ kJ}) = -1,300 \text{ kJ}$.

17. **D**

Choice (D) is the only reaction where the number of moles of gas is increasing, going from 2 moles of gas on the reactant side to 3 moles of gas on the product side. In all the other choices, the number of moles of gas either decreases or remains constant.

18. **A**

In an endothermic reaction, heat is transferred into the reaction system.

19. **B**

The reaction that forms 2 moles of $\text{HF}(g)$ from its constituent elements has an enthalpy change of -540 kJ . The heat of formation is given by the reaction that forms 1 mole from these elements, so you can just divide -540 kJ by 2 to get -270 kJ .

20. **B**

For the temperature of the water to increase, the reaction must be exothermic. If this is the case, the hydration energy of the solution exceeds the lattice energy within the solute.

21. **C**

The reaction must be exothermic, as the water heats up. In an exothermic reaction, heat is transferred from the system (the reaction itself) to the surroundings (the water).

22. **C**

The reaction is exothermic, meaning it has a negative ΔH , which indicates that it is favored in terms of enthalpy. It is also favored in terms of entropy, as the ions start organized into the ionic lattice, and end up in a more disordered state as free-floating ions in solution.

23. **A**

Hydration energy (like lattice energy) is based on Coulomb's Law, which has factors of both charge and size. The charges are the same in both substances, but fluoride is smaller than chloride. As size is in the denominator of Coulomb's Law, a smaller size means there will be more hydration energy.

24. **A**

The temperature increase is indicative of energy being released, meaning the reaction is exothermic. The entropy (disorder) of the system is increasing as it moves from three gas molecules to five.

Free-Response

1. (a) Use the entropy values in the table.

$$\Delta S^{\circ} = \sum S^{\circ}_{\text{products}} - \sum S^{\circ}_{\text{reactants}}$$

$$\Delta S^{\circ} = [(6)(213.6) + (6)(69.9)] - [(212.13) + (6)(205)]$$

J/mol×K

$$\Delta S^{\circ} = 259 \text{ J/mol} \times \text{K}$$

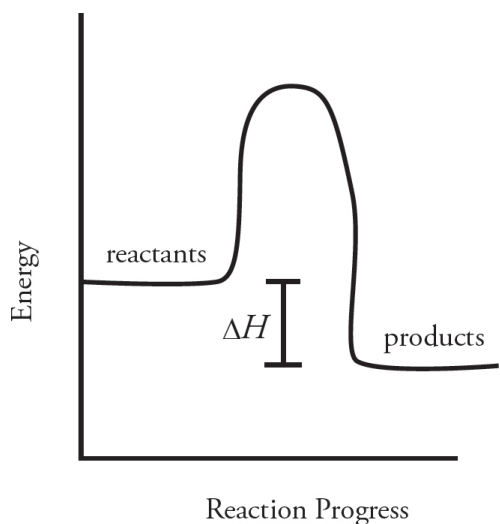
- (b) Use the equation below. Remember that enthalpy values are given in kJ and entropy values are given in J.

$$\Delta G^{\circ} = \Delta H^{\circ} - T\Delta S^{\circ}$$

$$\Delta G^{\circ} = (-2,801 \text{ kJ/mol}_{\text{rxn}}) - (298 \text{ K})(0.259 \text{ kJ/mol}_{\text{rxn}} \cdot \text{K}) =$$

-2,880 kJ/mol_{rxn}

(c)



The reaction is exothermic; therefore the reactants must have more energy than the products, as indicated. The difference in energy is equal to the ΔH for this reaction.

$$\begin{aligned} \text{(d) } 1.00 \text{ g glucose} &\times \frac{1 \text{ mol glucose}}{180 \text{ g glucose}} \times \\ &\frac{1 \text{ mol}_{\text{rxn}}}{1 \text{ mol glucose}} \times \frac{-2801 \text{ kJ}}{1 \text{ mol}_{\text{rxn}}} = -15.5 \text{ kJ} \end{aligned}$$

2. (a) Use the relationship below.

$$\Delta H^\circ = \sum \text{Energies of the bonds broken} - \sum \text{Energies of the bonds formed}$$

$$\Delta H^\circ = [(4)(415) + (2)(495)] - [(2)(799) + (4)(463)] \text{ kJ/mol}$$

$$\Delta H^\circ = -800 \text{ kJ/mol}$$

$$\begin{aligned} \text{(b) } 1,500 \text{ kJ} &\times \frac{1 \text{ mol CH}_4}{800 \text{ kJ}} = 1.9 \text{ mol} \\ \text{CH}_4 &\times \frac{16.04 \text{ g CH}_4}{1 \text{ mol CH}_4} = 30 \text{ g CH}_4 \end{aligned}$$

(c) Use $\Delta G^\circ = \Delta H^\circ - T\Delta S^\circ$

Remember that enthalpy values are given in kJ and entropy values are given in J.

$$\Delta S^\circ = \frac{\Delta H - \Delta G}{T} = \frac{(-800 \text{ kJ/mol}) - (-801 \text{ kJ/mol})}{(298 \text{ K})}$$

$$\Delta S^\circ = 0.003 \text{ kJ/K} = 3 \text{ J/K}$$

(d) ΔS° is very small, which means that the entropy change for the process is very small. This makes sense because the number of moles remains constant, the number of moles of gas remains constant, and the complexity of the molecules remains about the same.

3. (a) The heat of fusion is represented by the lower horizontal line, and the heat of vaporization by the higher one. As the higher line is longer, that means more heat is required to vaporize the substance than to melt it.
- (b) The substance is a gas at its highest temperature, and it would not change phase again.
- (c) The student is incorrect. The greater the specific heat of a substance, the more heat that is required to change the temperature of that substance. On the graph, that is represented by a shallower slope. As the slope of the solid (leftmost) line is greater than the liquid (central) or gas (rightmost) lines, the solid would actually have a lower specific heat than either the liquid or the gas.
- (d) Heat is being added to the system, so enthalpy change is positive. As the substance first melts, and then boils, it is becoming more disordered. Even after it becomes a gas, the gas molecules will continue to spread out as the

temperature increases. So, the entropy change is also positive.

4. (a) (i) In order to boil a liquid, the intermolecular forces between the various molecules of the liquid must be broken. This requires the input of energy, thus ΔH is positive.

(ii) A liquid turning into a gas leads to an increase in disorder, which yields a positive ΔS .

(b) When a phase change is occurring, the value for ΔG is always zero, as no real chemical reaction is occurring. Using that knowledge and making sure the units are matching (by converting the entropy to $\text{kJ/mol} \times \text{K}$), you get:

$$\Delta G = \Delta H^\circ - T\Delta S^\circ$$

$$0 = 37.6 \text{ kJ/mol} - T(0.111 \text{ kJ/mol} \times \text{K})$$

$$T = 339 \text{ K}$$

$$\text{Converting to Celsius: } 339 \text{ K} - 273.15 = 65.9^\circ\text{C}$$

(c) First, figure out how many moles of ethanol are present.

$$D = \frac{m}{V} \quad 0.789 \text{ g/mL}$$
$$= \frac{m}{50.0 \text{ mL}} \quad m = 39.5 \text{ g CH}_3\text{OH}$$

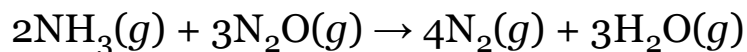
$$39.5 \text{ g} \times \frac{1 \text{ mol CH}_3\text{OH}}{32.05 \text{ g}} = 1.23 \text{ mol CH}_3\text{OH}$$

Then, use the ΔH° value to calculate the amount of heat needed.

$$1.23 \text{ mol} \times \frac{37.6 \text{ kJ}}{\text{mol}} = 46.2 \text{ kJ}$$

- (d) As methanol (or any other liquid) boils, the temperature will remain constant, as the added heat is going into breaking intermolecular forces instead of adding extra kinetic energy to the liquid.
- (e) Yes, methanol would be soluble with water. Methanol is a polar molecule, as is water, and in terms of molecular solubility, like dissolves like.
- (f) It would be greater than that of methanol. Both molecules are polar with H-bonds, but ethanol has more electrons, and thus its electron cloud is more polarizable. This causes the London dispersion forces in ethanol to be stronger than those in methanol, leading to a higher boiling point.

5. Ammonia gas reacts with dinitrogen monoxide via the following reaction:



The standard entropy of formation values for the varying substances are listed in the table below.

Substance	S° (J/mol·K)
$\text{NH}_3(g)$	193
$\text{N}_2\text{O}(g)$	220
$\text{N}_2(g)$	192
$\text{H}_2\text{O}(g)$	189

- (a) Calculate the entropy value for the overall reaction.

$$\Delta S = \Delta S_{\text{products}} - \Delta S_{\text{reactants}}$$

$$\Delta S = [3(\text{H}_2\text{O}) + 4(\text{N}_2)] - [2(\text{NH}_3) + 3(\text{N}_2\text{O})]$$

$$\Delta S = [3(189) + 4(192)] - [2(193) + 3(220)]$$

$$\Delta S = 1335 - 1046$$

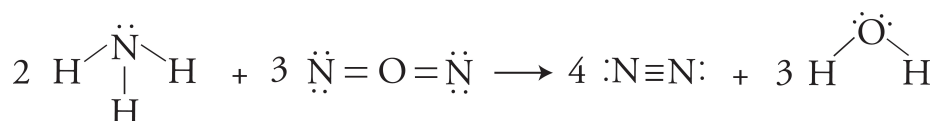
$$\Delta S = 289 \text{ J/mol}\cdot\text{K}$$

Several bond enthalpies are listed in the table below.

Bond	Enthalpy (kJ/mol)	Bond	Enthalpy (kJ/mol)
N–H	388	N=N	409
N–O	210	N≡N	941
N=O	630	O–H	463

(b) Calculate the enthalpy value for the overall reaction.

To determine the amount of enthalpy change, Lewis structures of all of the species should be drawn.



You have to account for not only how many bonds are within each molecule, but also how many of those molecules there are. For instance, there are three N–H bonds in NH_3 , so with two NH_3 molecules there will be six total N–H bonds broken. The bond energy for the other three molecules must be calculated the same way.

$$\Delta H = \text{Bonds broken (reactants)} - \text{bonds formed (products)}$$

$$\Delta H = [6(\text{N-H}) + 6(\text{N=O})] - [4(\text{N}\equiv\text{N}) + 6(\text{O-H})]$$

$$\Delta H = [6(388) + 6(630)] - [4(941) + 6(463)]$$

$$\Delta H = 6108 - 6542$$

$$\Delta H = -434 \text{ kJ/mol}$$

(c) Is this reaction thermodynamically favored at 25°C? Justify your answer.

We will use the Gibbs free energy equation here. Keep in mind the units have to match. In this solution, the units for entropy have been converted to kJ.

$$\Delta G = \Delta H - T\Delta S$$

$$\Delta G = -434 \text{ kJ/mol} - (298 \text{ K})(0.289 \text{ kJ/mol}\times\text{K})$$

$$\Delta G = -434 \text{ J/mol} - 86.1 \text{ kJ/mol}$$

$$\Delta G = -520 \text{ kJ/mol}$$

The value for ΔG is negative, therefore the reaction is thermodynamically favored at 25°C.

(d) If 25.00 g of NH_3 reacts with 25.00 g of N_2O :

(i) Will energy be released or absorbed?

The value for ΔH is negative; therefore it is an exothermic reaction and energy will be released.

(ii) What is the magnitude of the energy change?

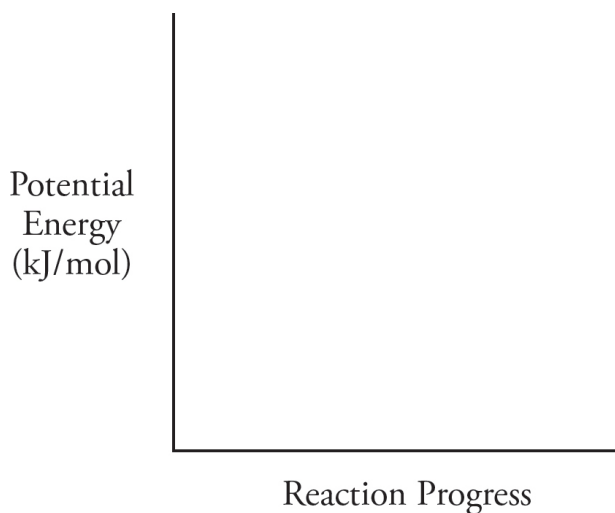
To determine how much energy is released, the limiting reagent must be determined.

$$25.00 \text{ g NH}_3 \times \frac{1 \text{ mol NH}_3}{17.04 \text{ g NH}_3} \times \frac{1 \text{ mol}_{\text{rxn}}}{2 \text{ mol NH}_3} \times \frac{-434 \text{ kJ}}{1 \text{ mol}_{\text{rxn}}} = -318.4 \text{ kJ}$$

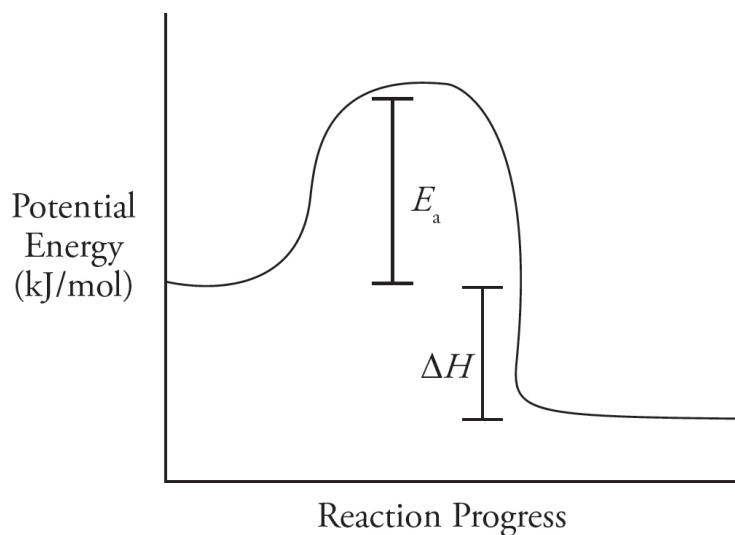
$$25.00 \text{ g N}_2\text{O} \times \frac{1 \text{ mol N}_2\text{O}}{44.02 \text{ g N}_2\text{O}} \times \frac{1 \text{ mol}_{\text{rxn}}}{3 \text{ mol N}_2\text{O}} \times \frac{-434 \text{ kJ}}{1 \text{ mol}_{\text{rxn}}} = -82.16 \text{ kJ}$$

As the N_2O would produce less energy, it would run out first and is thus limiting. The answer is thus -82.16 kJ .

- (e) On the reaction coordinates below, draw a line showing the progression of this reaction. Label both ΔH and E_a on the graph.



The activation energy describes the distance between the bond energy of the reactants and the energy of the activated state. In terms of enthalpy, the value for ΔH is negative, so that means that the bond energy of the products will be lower than that of the reactants.



6. (a) $q = mc\Delta T$

$$q = (25.0 \text{ g})(4.18 \text{ J/g}\times^{\circ}\text{C})(3.6^{\circ}\text{C})$$

$$q = 376 \text{ J}$$

(b) The heat gained by the water is the same as the heat lost by the aluminum.

$$q = mc\Delta T$$

$$-376 \text{ J} = (5.86 \text{ g})(c)(-82.3^{\circ}\text{C})$$

$$c = 0.780 \text{ J/g}\times^{\circ}\text{C}$$

$$(c) 5.86 \text{ g Al} \times \frac{1 \text{ mol Al}}{26.98 \text{ g Al}} = 0.217 \text{ mol Al}$$

$$376 \text{ J} / 0.217 \text{ mol} = 1730 \text{ J/mol} = 1.73 \text{ kJ/mol}$$

$$(d) \% \text{ error} = \frac{|\text{experimental} - \text{accepted}|}{\text{accepted}} \times 100\%$$

$$\frac{|0.780 - 0.900|}{0.900} \times 100\% = 13.3\% \text{ error}$$

(e) Error 1: If some of the heat that was lost by the aluminum was not absorbed by the water, that would cause the calculated heat gained by the water in part (a) to be artificially low. This, in turn, would reduce the value of the specific heat of aluminum as calculated in part (b).

Error 2: If there was more than 25.0 mL of water in the calorimeter, that would mean the mass in part (a) was artificially low, which would make the calculation for the heat gained by the water also artificially low. This, in turn, would reduce the value of the specific heat of aluminum calculated in part (b).

There are many potential errors here, but as long as you can quantitatively follow them to the conclusion that the experimental value would be too low, any error (within reason) can be acceptable.

Chapter 8

Big Idea #6: Equilibrium, Acids and Bases, Titrations, and Solubility

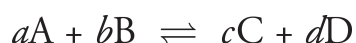
Any bond or intermolecular attraction that can be formed can be broken. These two processes are in dynamic competition, sensitive to initial conditions and external perturbations.

THE EQUILIBRIUM CONSTANT, K_{eq}

Most chemical processes are reversible. That is, reactants react to form products, but those products can also react to form reactants. A reaction is at equilibrium when the rate of the forward reaction is equal to the rate of the reverse reaction. The relationship between the concentrations of reactants and products in a reaction at equilibrium is given by the equilibrium expression, also called the **Law of Mass Action**.

The Equilibrium Expression

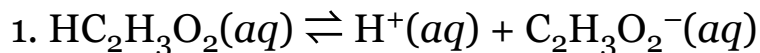
For the reaction



$$K_{eq} = \frac{[C]^c [D]^d}{[A]^a [B]^b}$$

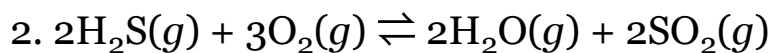
1. $[A]$, $[B]$, $[C]$, and $[D]$ are molar concentrations or partial pressures at equilibrium.
2. Products are in the numerator, and reactants are in the denominator.
3. Coefficients in the balanced equation become exponents in the equilibrium expression.
4. Solids and pure liquids are not included in the equilibrium expression because they cannot change their concentration. Only gaseous and aqueous species are included in the expression.
5. Units are not given for K_{eq} .

Let's look at a few examples:



$$K_{eq} = K_a = \frac{[\text{H}^+][\text{C}_2\text{H}_3\text{O}_2^-]}{[\text{HC}_2\text{H}_3\text{O}_2]}$$

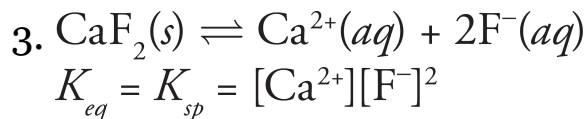
This reaction shows the dissociation of acetic acid in water. All of the reactants and products are aqueous particles, so they are all included in the equilibrium expression. None of the reactants or products have coefficients, so there are no exponents in the equilibrium expression. This is the standard form of K_a , the acid dissociation constant.



$$K_{eq} = K_c = \frac{[\text{H}_2\text{O}]^2 [\text{SO}_2]^2}{[\text{H}_2\text{S}]^2 [\text{O}_2]^3}$$

$$K_{eq} = K_p = \frac{(P_{\text{H}_2\text{O}})^2 (P_{\text{SO}_2})^2}{(P_{\text{H}_2\text{S}})^2 (P_{\text{O}_2})^3}$$

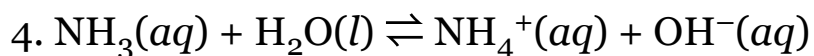
All of the reactants and products in this reaction are gases, so K_{eq} can be expressed in terms of concentration (K_c , moles/liter or molarity) or in terms of partial pressure (K_p , atmospheres). All of the reactants and products are included here, and the coefficients in the reaction become exponents in the equilibrium expression.



$$K_{eq} = K_{sp} = [\text{Ca}^{2+}][\text{F}^-]^2$$

This reaction shows the dissociation of a slightly soluble salt. There is no denominator in this equilibrium expression because the reactant

is a solid. Solids are left out of the equilibrium expression because the concentration of a solid is constant. There must be some solid present for equilibrium to exist, but you do not need to include it in your calculations. This form of K_{eq} is called the solubility product, K_{sp} .



$$K_{eq} = K_b = \frac{[\text{NH}_4^+][\text{OH}^-]}{[\text{NH}_3]}$$

This is the acid-base reaction between ammonia and water. We can leave water out of the equilibrium expression because it is a pure liquid. This is the standard form for K_b , the base dissociation constant.

Here is a roundup of the equilibrium constants you need to be familiar with for the test.

- K_c is the constant for molar concentrations.
- K_p is the constant for partial pressures.
- K_{sp} is the solubility product, which has no denominator because the reactants are solids.
- K_a is the acid dissociation constant for weak acids.
- K_b is the base dissociation constant for weak bases.
- K_w describes the ionization of water ($K_w = 1 \times 10^{-14}$).

The equilibrium constant has a lot of aliases, but they all take the same form and tell you the same thing. The **equilibrium constant** tells you the relative amounts of products and reactants at equilibrium.

A large value for K_{eq} means that products are favored over reactants at equilibrium, while a small value for K_{eq} means that reactants are favored over products at equilibrium.

Manipulating K_{eq}

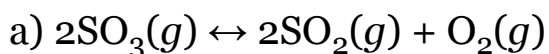
Much as Hess's Law allows for the determination of the enthalpy for a reaction given the enthalpy values for similar reactions, you can determine the equilibrium constant of a reaction by manipulating similar reactions with known equilibrium constants. However, the rules for doing so are different than the rules for shifting enthalpy values:

1. If you flip a reaction, you take the reciprocal of the equilibrium constant to get the new equilibrium constant.
2. If you multiply a reaction by a coefficient, you take the equilibrium constant to that power to get the new constant.
3. If you add two reactions together, you multiply the equilibrium constants of those reactions to get the new constant.

Example:

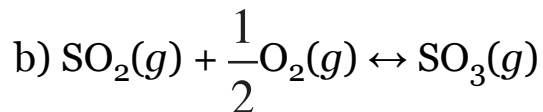


According to the information above, what is the equilibrium constant for each of the following reactions?



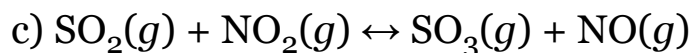
This reaction is the opposite of the first reaction, so

$$K = \frac{1}{4.0 \times 10^{24}} = 2.5 \times 10^{-25}.$$

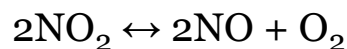


This reaction is the first reaction multiplied by one-half. Thus, the new equilibrium constant is:

$$K = (4.0 \times 10^{24})^{1/2} = 2.0 \times 10^{12}$$

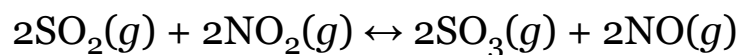


To get this reaction, we first have to flip the second reaction:



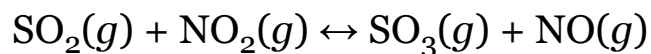
$$K = \frac{1}{0.065} = 16$$

We can then add the flipped second reaction to the original first reaction. The O_2 will cancel out, yielding the following:



$$K = 4.0 \times 10^{24}(16) = 6.4 \times 10^{25}$$

Finally, we have to multiply that equation by half to get our desired reaction.



$$K = (6.4 \times 10^{25})^{1/2} = 2.5 \times 10^{12}$$

LE CHÂTELIER'S PRINCIPLE

At equilibrium, the rates of the forward and reverse reactions are equal. A “shift” in a certain direction means the rate of the forward or reverse reaction increases. Le Châtelier’s Principle states that whenever a stress is placed on a system at equilibrium, the system will shift in response to that stress. If the forward rate increases, we say the reaction has shifted right, which will create more products. If the reverse rate increases, we say the reaction has shifted left, which will create more reactants.

Let’s use the **Haber process**, which is used in the industrial preparation of ammonia, as an example.



Concentration

When the concentration of a reactant or product is increased, the reaction will shift in the direction that allows it to use up the added substance. If N_2 or H_2 is added, the reaction shifts right. If NH_3 is added, the reaction shifts left.

When the concentration of a species is decreased, the reaction will shift in the direction that allows it to create the substance that has been removed. If N_2 or H_2 is removed, the reaction shifts left. If NH_3 is removed, the reaction shifts right.

Pressure

When the external pressure on a system is increased, that will increase the partial pressures of all of the gases inside the container and cause a shift to the side with less gas molecules. In the Haber process, there are four gas molecules on the left and two on the right, so an increase in partial pressures would cause a shift to the right. Decreasing the partial pressure of all gases would have the opposite effect and cause a shift to the left.

The most common way to cause a change in pressure is by changing the size of the container in which a reaction is occurring. If the Haber process were occurring inside a balloon, and we were to decrease the total volume of that balloon by squeezing it, that would increase the partial pressures of all gases involved (assuming the temperature stayed constant). Increasing the volume of the container would have the opposite effect.

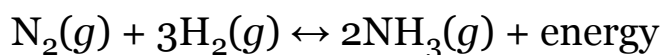


In addition to changing volume, the partial pressure of gases can be changed by adding another gas to the container while maintaining the total pressure, even if that gas does not participate in the reaction. If we were to add some helium gas to a container where the Haber process was in equilibrium, it would decrease the mole fraction of all of the gases participating in the reaction (since the helium now represents at least some percentage of the total gases). A decreased mole fraction causes a decreased partial pressure, which again causes a shift to the left.

Note that for any equilibrium, there has to be a different number of gas molecules on each side of the equilibrium for a pressure change to cause any kind of shift. In equilibria where the number of gas molecules are the same on both sides, any of the above changes would cause no shift at all.

Temperature

There's a trick to figure out what happens when the temperature changes. First, rewrite the equation to include the heat energy on the side that it would be present on. The Haber process is exothermic, so heat is generated:



Then, if the temperature goes up, the reaction will proceed in the reverse direction (shifting away from the added energy). If the

temperature goes down, the reaction will proceed in the forward direction (creating more energy). The reverse would be true in an endothermic reaction, as the energy would be part of the reactants.

Dilution

One last type of shift which is caused by dilution. For an aqueous equilibrium, diluting that equilibrium with extra water can cause a shift. The Haber process won't work for this one, so let's examine another equilibrium:



If you were to dilute the above equilibrium with extra water, it would shift to the side that has more aqueous species. In the above example, this causes a shift to the left. If some water were removed (say, by evaporation), the reaction would shift to the side with less aqueous species (right).

CHANGES IN THE EQUILIBRIUM CONSTANT

It is important to understand that shifts caused by concentration or pressure changes are temporary shifts, and do not change the value of the equilibrium constant itself. Eventually, the concentrations of the products and reactants will re-establish the same ratio as they originally had at equilibrium.

However, shifts caused by temperature changes are different. Because changing temperature also affects reaction kinetics by adding (or removing) energy from the equilibrium system, a change in temperature will also affect the equilibrium constant for the reaction itself, in addition to causing a shift. This means the ratio of the products to reactants at equilibrium will change as the temperature changes.

In the Haber process, an increase in temperature causes a shift to the left, and it would permanently affect the value of the equilibrium

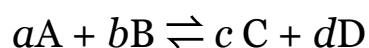
constant in a way that is consistent with the shift. A shift to the left causes an increase in the concentrations of the reactants (the denominator in the mass action expression) while simultaneously causing a decrease in the concentration of the product (the numerator in the mass action expression). Thus, increasing the temperature decreases the value of the equilibrium constant—that is, it causes there to be a larger amount of reactants present at equilibrium compared to the amount of products present. The reverse would be true for a temperature decrease; the shift to the right would cause an increase in the equilibrium constant.

THE REACTION QUOTIENT, Q

The reaction quotient is essentially the quantitative application of Le Châtelier's Principle. It is determined using the Law of Mass Action. However, you can use the concentration or pressure values at any point in the reaction to calculate Q . (Remember, you can only use concentrations or pressures of reactions at equilibrium to calculate K). The value for the reaction quotient can be compared to the value for the equilibrium constant to predict in which direction a reaction will shift from the given set of initial conditions.

The Reaction Quotient

For the reaction



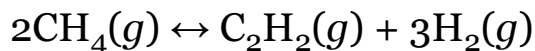
$$Q = \frac{[C]^c [D]^d}{[A]^a [B]^b}$$

$[A]$, $[B]$, $[C]$, and $[D]$ are initial molar concentrations or partial pressures.

- If Q is less than K , the reaction shifts right.

- If Q is greater than K , the reaction shifts left.
- If $Q = K$, the reaction is already at equilibrium.

Let's take a look at an example. The following reaction takes place in a sealed flask:



(a) Determine the equilibrium constant if the following concentrations are found at equilibrium.

$$[\text{CH}_4] = 0.0032 \text{ M} \quad [\text{C}_2\text{H}_2] = 0.025 \text{ M} \quad [\text{H}_2] = 0.040 \text{ M}$$

To solve this, we first need to create the equilibrium constant expression:

$$K_c = \frac{[\text{C}_2\text{H}_2][\text{H}_2]^3}{[\text{CH}_4]^2}$$

By plugging in the numbers, we get:

$$K_c = \frac{(0.025)(0.040)^3}{(0.0032)^2} \quad K_c = \frac{1.6 \times 10^{-6}}{1.0 \times 10^{-5}} \quad K_c = 0.16$$

(b) Upon testing this reaction at another point, the following concentrations are found:

$$[\text{CH}_4] = 0.0055 \text{ M} \quad [\text{C}_2\text{H}_2] = 0.026 \text{ M} \quad [\text{H}_2] = 0.029 \text{ M}$$

Use the reaction quotient to determine which way the reaction needs to shift to reach equilibrium.

Q uses the exact same ratios as the equilibrium constant expression, so:

$$Q = \frac{[\text{C}_2\text{H}_2][\text{H}_2]^3}{[\text{CH}_4]^2} = \frac{(0.026)(0.029)^3}{(0.0055)^2} = \frac{6.3 \times 10^{-7}}{3.0 \times 10^{-5}} = 0.021$$

0.021 is less than the equilibrium constant value of 0.16, so the reaction must proceed to the right, creating more products and reducing the amount of reactant in order to come to equilibrium.

Voltaic Cells

When examining voltaic cells, the reduction potentials are always given at standard conditions; that is 25°C, 1.0 atm, and with all species having a concentration of 1.0 *M*. If any of those conditions deviate, it will also cause the cell potential to deviate.

Voltaic cells are all very favored, having equilibrium constants significantly greater than 1. If the reaction quotient for a voltaic cell were to ever become equal to the equilibrium constant (so, at $Q = K$), the voltage of the cell would drop to zero. Given that knowledge, the best way to determine how a cell's potential will change if standard conditions are deviated from is to use the reaction quotient.

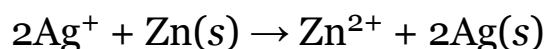
As all concentrations are equal to 1.0 *M* at standard conditions, we can infer that at standard conditions, the reaction quotient will also be equal to 1. Any change to the cell that would cause the reaction quotient to increase (say, an increase in the concentration of a product or the decrease in the concentration of a reactant) would cause the reaction quotient to become closer to the equilibrium constant and would thus decrease the cell potential (remember, if Q ever reaches K , the cell potential becomes zero!). Any change that would cause the reaction quotient to decrease would move it further from the equilibrium constant, and thus increase the potential of the cell.

If a cell has a gas at either the cathode or the anode, a change in pressure can also affect the reaction quotient, and a change in temperature may as well. However, concentration changes, which

usually occur constantly as the reaction in a cell is progressing, are the most common reason that a voltaic cell will deviate from its standard potential.

Example:

A voltaic cell using silver and zinc is connected, and the below reaction occurs.



What would happen to the voltage of the above cell as the reaction progresses?

As the reaction progresses, $[\text{Ag}^+]$ decreases as it is reduced into Ag(s) , and $[\text{Zn}^{2+}]$ increases as it is oxidized from Zn(s) . Both of these changes would have the effect of increasing the reaction quotient, which would bring the cell closer to equilibrium and thus decrease the overall potential.

SOLUBILITY

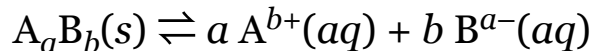
Roughly speaking, a salt can be considered “soluble” if more than 1 gram of the salt can be dissolved in 100 milliliters of water. Soluble salts are usually assumed to dissociate completely in aqueous solution. Most, but not all, solids become more soluble in a liquid as the temperature is increased.

Solubility Product (K_{sp})

Salts that are “slightly soluble” and “insoluble” still dissociate in solution to some extent. The solubility product (K_{sp}) is a measure of the extent of a salt’s dissociation in solution. The K_{sp} is one of the forms of the equilibrium expression. The greater the value of the solubility product for a salt, the more soluble the salt.

Solubility Product

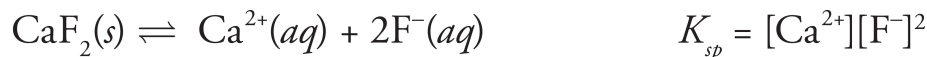
For the reaction



The solubility expression is

$$K_{sp} = [A^{b+}]^a [B^{a-}]^b$$

Here are some examples:



The solubility of salts can be described by the K_{sp} or by the molar solubility. The molar solubility of the salt describes the number of moles of salt that can be dissolved per liter of solution. The K_{sp} value for Ag_2CrO_4 is 8.0×10^{-12} . We can determine the molar solubility using the following calculations:

$$K_{sp} = [Ag^{+}]^2[CrO_4^{2-}]$$

$$8.0 \times 10^{-12} = (2x)^2(x)$$

$$8.0 \times 10^{-12} = 4x^3$$

$$x = [CrO_4^{2-}] = 1.3 \times 10^{-4} M$$

$$2x = [Ag^{+}] = 2.6 \times 10^{-4} M$$

The molar solubility of the salt will also be equal to the concentration of any ion that occurs in a 1:1 ratio with the salt. In the example above, the molar solubility of the salt is $1.3 \times 10^{-4} M$, the same as the concentration of the CrO_4^{2-} ions.

Note that the concentration of the silver ions is both doubled (because twice as many of them will be in solution) AND squared in the K_{sp} expression. Thus, the 2 coefficient on the silver is represented twice in the solubility product calculations. Forgetting to account for that coefficient in the ion concentration (like using just x instead of $2x$) is a very common mistake made by students; take care to avoid it!

Typically, the molar solubility of most salts will increase with rising temperatures. This is because a higher temperature has more energy available to force the water molecules apart and make room for the solute ions. There are some salts that see their solubility decrease with increasing temperature, but there is no easy way to predict when that will occur.

The Common Ion Effect

Let's look at the solubility expression for AgCl.

$$K_{sp} = [\text{Ag}^+][\text{Cl}^-] = 1.6 \times 10^{-10}$$

If we throw a block of solid AgCl into a beaker of water, we can tell from the K_{sp} what the concentrations of Ag^+ and Cl^- will be at equilibrium. For every unit of AgCl that dissociates, we get one Ag^+ and one Cl^- , so we can solve the equation above as follows:

$$[\text{Ag}^+][\text{Cl}^-] = 1.6 \times 10^{-10}$$

$$(x)(x) = 1.6 \times 10^{-10}$$

$$x^2 = 1.6 \times 10^{-10}$$

$$x = [\text{Ag}^+] = [\text{Cl}^-] = 1.3 \times 10^{-5} M$$

So there are very small amounts of Ag^+ and Cl^- in the solution.

Let's say we add 0.10 mole of NaCl to 1 liter of the AgCl solution. NaCl dissociates completely, so that's the same thing as adding 0.1 mole of Na^+ ions and 0.1 mole of Cl^- ions to the solution. The Na^+ ions will not affect the AgCl equilibrium, so we can ignore them; but the Cl^- ions must be taken into account. That's because of the **common ion effect**.

The common ion effect says that the newly added Cl^- ions will affect the AgCl equilibrium, although the newly added Cl^- ions did not come from AgCl.

Let's look at the solubility expression again. Now we have 0.10 mole of Cl^- ions in 1 liter of the solution, so $[\text{Cl}^-] = 0.10 M$.

$$[\text{Ag}^+][\text{Cl}^-] = 1.6 \times 10^{-10}$$

$$[\text{Ag}^+](0.10 M) = 1.6 \times 10^{-10}$$

$$[\text{Ag}^+] = \frac{(1.6 \times 10^{-10})}{(0.10)} M$$

$$[\text{Ag}^+] = 1.6 \times 10^{-9} M$$

Now the number of Ag^+ ions in the solution has decreased drastically because of the Cl^- ions introduced to the solution by NaCl. So when solutions of AgCl and NaCl, which share a common Cl^- ion, are

mixed, the more soluble salt (NaCl) can cause the less soluble salt (AgCl) to precipitate. In general, when two salt solutions that share a common ion are mixed, the salt with the lower value for K_{sp} will precipitate first.

Standard Free Energy Change and the Equilibrium Constant

The amount of Gibbs free energy in any given reaction can also be calculated if you know the equilibrium constant expression for that reaction.

$$\Delta G^\circ = -RT \ln K$$

R = the gas constant, 8.31 J/mol $\times$ K

T = absolute temperature (K)

K = the equilibrium constant

Note that the “ R ” value here is NOT the same as the “ R ” value that we use when dealing with ideal gases! If ΔG° is negative, K must be greater than 1, and products will be favored at equilibrium. Alternatively, if ΔG° is positive, K must be less than 1, and reactants will be favored at equilibrium.

ACIDS AND BASES DEFINITIONS

Arrhenius

S. A. Arrhenius defined an acid as a substance that ionizes in water and produces hydrogen ions (H^+). As hydrogen atoms consist of one single proton and one single electron, removing that electron when we form the ion leaves just the proton behind. For that reason, you

will often hear H^+ ions referred to simply as protons. For instance, HCl is an acid.



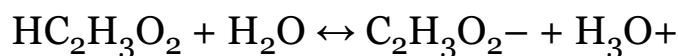
He defined a base as a substance that ionizes in water and produces hydroxide ions (OH^- ions). For instance, NaOH is a base.



Brønsted-Lowry

J. N. Brønsted and T. M. Lowry defined an acid as a substance that is capable of donating a proton, which is the same as donating an H^+ ion, and they defined a base as a substance that is capable of accepting a proton. This definition is the one that will be used most frequently on the exam.

Look at the reversible reaction below.

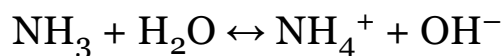


According to Brønsted-Lowry

$\text{HC}_2\text{H}_3\text{O}_2$ and H_3O^+ are acids.

$\text{C}_2\text{H}_3\text{O}_2^-$ and H_2O are bases.

Now look at this reversible reaction.

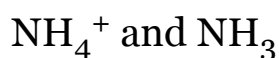


According to Brønsted-Lowry

NH_3 and OH^- are bases.

H_2O and NH_4^+ are acids.

So in each case, the species with the H^+ ion is the acid, and the same species without the H^+ ion is the base; the two species are called a **conjugate pair**. The following are the acid-base conjugate pairs in the reactions above:



Notice that water can act either as an acid or base. Any substance which has that ability is called **amphoteric**.

pH

Many of the concentration measurements in acid-base problems are given to us in terms of pH and pOH.

$$\text{p (anything)} = -\log (\text{anything})$$

$$\text{pH} = -\log [\text{H}^+]$$

$$\text{pOH} = -\log [\text{OH}^-]$$

$$\text{p}K_a = -\log K_a$$

$$\text{p}K_b = -\log K_b$$

In a solution

- when $[\text{H}^+] = [\text{OH}^-]$, the solution is neutral, and $\text{pH} = 7$

- when $[H^+]$ is greater than $[OH^-]$, the solution is acidic, and pH is less than 7
- when $[H^+]$ is less than $[OH^-]$, the solution is basic, and pH is greater than 7

It is important to remember that *increasing* pH means *decreasing* $[H^+]$, which means that there are fewer H^+ ions floating around and the solution is *less acidic*. Alternatively, *decreasing* pH means *increasing* $[H^+]$, which means that there are more H^+ ions floating around and the solution is *more acidic*.

ACID STRENGTHS

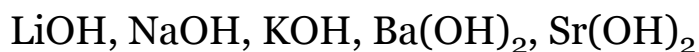
Strong Acids

Strong acids dissociate completely in water, so the reaction goes to completion and they never reach equilibrium with their conjugate bases. Because there is no equilibrium, there is no equilibrium constant, so there is no dissociation constant for strong acids or bases.

Important Strong Acids



Important Strong Bases



Because the dissociation of a strong acid goes to completion, there is no tendency for the reverse reaction to occur, which means that the conjugate base of a strong acid must be extremely weak.

It's much easier to find the pH of a strong acid solution than it is to find the pH of a weak acid solution. That's because strong acids dissociate completely, so the final concentration of H^+ ions will be the same as the initial concentration of the strong acid.

Let's look at a 0.010-molar solution of HCl.

HCl dissociates completely, so $[\text{H}^+] = 0.010\text{ M}$

$$\text{pH} = -\log [\text{H}^+] = -\log (0.010) = -\log (10^{-2}) = 2$$

So you can always find the pH of a strong acid solution directly from its concentration.

Weak Acids

When a weak acid (often symbolized with HA) is placed in water, a small fraction of its molecules will dissociate into hydrogen ions (H^+) and conjugate base ions (A^-). Most of the acid molecules will remain in solution as undissociated aqueous particles.

The dissociation constants, K_a and K_b , are measures of the strengths of weak acids and bases. K_a and K_b are just the equilibrium constants specific to acids and bases.

Acid Dissociation Constant

$$K_a = \frac{[\text{H}^+][\text{A}^-]}{[\text{HA}]}$$

$[\text{H}^+]$ = concentration of hydrogen ions (M)

$[\text{A}^-]$ = concentration of conjugate base ions (M)

$[\text{HA}]$ = concentration of undissociated acid molecules (M)

Base Dissociation Constant

$$K_b = \frac{[\text{HB}^+][\text{OH}^-]}{[\text{B}]}$$

$[\text{HB}^+]$ = concentration of conjugate acid ions (M)

$[\text{OH}^-]$ = concentration of hydroxide ions (M)

$[\text{B}]$ = concentration of unprotonated base molecules (M)

The greater the value of K_a , the greater the extent of the dissociation of the acid and the stronger the acid. The same thing goes for K_b , but in the case of K_b , the base is not dissociating. Instead, it is accepting a hydrogen ion (a proton) from an acid. So, a base does not dissociate—it protonates (or ionizes).

If you know the K_a for an acid and the concentration of the acid, you can find the pH. For instance, let's look at 0.20-molar solution of $\text{HC}_2\text{H}_3\text{O}_2$, with $K_a = 1.8 \times 10^{-5}$.

First, we set up the K_a equation, plugging in values.



$$K_a = \frac{[\text{H}^+][\text{C}_2\text{H}_3\text{O}_2^-]}{[\text{HC}_2\text{H}_3\text{O}_2]}$$

Therefore, the ICE (Initial, Change, Equilibrium) table for the above problem is as follows:

	$[\text{HC}_2\text{H}_3\text{O}_2]$	$[\text{H}^+]$	$[\text{C}_2\text{H}_3\text{O}_2^-]$

Initial	0.20	0.0	0.0
Change	$-x$	$+x$	$+x$
Equilibrium	$0.20 - x$	x	x

Because every acid molecule that dissociates produces one H^+ and one $\text{C}_2\text{H}_3\text{O}_2^-$,

$$[\text{H}^+] = [\text{C}_2\text{H}_3\text{O}_2^-] = x$$

and because, strictly speaking, the molecules that dissociate should be subtracted from the initial concentration of $\text{HC}_2\text{H}_3\text{O}_2$, $[\text{HC}_2\text{H}_3\text{O}_2]$ should be $(0.20\text{ M} - x)$. In practice, however, x is almost always insignificant compared with the initial concentration of acid, so we just use the initial concentration in the calculation.

$$[\text{HC}_2\text{H}_3\text{O}_2] = 0.20\text{ M}$$

Now we can plug our values and variable into the K_a expression.

$$1.8 \times 10^{-5} = \frac{x^2}{0.20}$$

Solve for x .

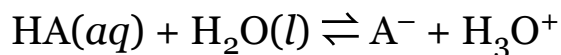
$$x = [\text{H}^+] = 1.9 \times 10^{-3}$$

Now that we know $[\text{H}^+]$, we can calculate the pH.

$$\text{pH} = -\log [\text{H}^+] = -\log (1.9 \times 10^{-3}) = 2.7$$

This is the basic approach to solving many of the weak acid/base problems that will be on the test.

Another way to write out the dissociation of a weak acid is:



The above includes water molecules in the reaction, and it is technically more accurate overall, as an acid will not dissociate unless it has a base to give protons to. The H_3O^+ ion is called the hydronium ion, and you can substitute it for H^+ in any acid/base reaction. So, $-\log [\text{H}_3\text{O}^+] = \text{pH}$.

In solving the above problem, you were told that the “ $-x$ ” term in “ $0.20 - x$ ” (henceforth called the dissociation value) is almost always insignificant. In that problem, “ x ” turned out to be 0.0019, and if you plug that into the original term you get $0.20 - 0.0019 = 0.20$ (when using significant figures correctly). Those values are just approximations, of course, but the point is that the very small amount of acid that dissociates does not change the effective concentration of the acid.

That being said, if the weak acid has a high enough K_a value, the dissociation value can become significant. It is true that you will never be asked to solve a problem like the one above in which the dissociation value term is significant, because that would require using the quadratic equation, which is not required on the exam. However, if instead of being given the K_a and concentration values for a weak acid, you are given the pH and asked to solve for either the acid's concentration or K_a , you should not neglect the dissociation value. The best way to show this is through another example.

A 0.50 M solution of HClO has a pH of 1.13. Determine the K_a value for HClO .

First, let's look at the K_a expression.

$$K_a = \frac{[\text{H}^+][\text{ClO}^-]}{[\text{HClO}]}$$

The ICE chart would then look like this:

	[HClO]	[H ⁺]	[ClO ⁻]
Initial	0.50	0	0
Change	-x	+x	+x
Equilibrium	0.50 - x	x	x

As we are given the pH, we can easily calculate [H⁺], which is equal to x.

$$1.13 = -\log[\text{H}^+]$$

$$[\text{H}^+] = 0.074 \text{ M}$$

Plugging those numbers into the K_a expression yields:

$$K_a = \frac{(0.074)(0.074)}{(0.50 - 0.074)}$$

$$K_a = \frac{(0.074)(0.074)}{(0.43)}$$

$$K_a = 0.013 = 1.3 \times 10^{-2}$$

Solving without accounting for the dissociation value in the denominator here would have yielded a similar, but not identical, result. Even though the result would be similar, the points awarded on problems like these often focus on the process just as much as the actual result, which is why it is important to make sure the dissociation value is accounted for.

The exam is written in a way that is kind of tricky, in that it is safe to ignore the dissociation value when attempting to solve for pH because not doing so would lead to a quadratic equation, which is a

skill not required on the exam. However, it is often **not** safe to ignore the dissociation value when attempting to solve for K_a or acid concentration. The best advice, therefore, is to only ignore the dissociation value if using it would lead to a quadratic equation.

Percent Dissociation

The primary factor when it comes to determining acid strength is that the more H^+ ions that an acid can donate, the stronger that acid will be. How easily an acid dissociates is often determined in part by its structure.

Consider the binary acids composed of hydrogen and a halogen. Of those, HI, HBr, and HCl are all strong acids, meaning they dissociate completely. HF, however, is not a strong acid. The reason for this is that fluorine is extremely electronegative, and thus fluorine will “hold on” to the hydrogen more effectively.

If you consider oxoacids, though, the reverse trend is true. Consider the Lewis diagrams for HOF and HOBr:



In this case, fluorine’s very high electronegativity affects the O–F bond, drawing the shared electrons toward the fluorine. However, fluorine is so electronegative that it also attracts the shared electrons in the H–O bond as well, which weakens the overall H–O bond and makes the H more likely to dissociate. Thus, HOF is a stronger acid than HOBr.

The strength of an acid is very case-specific, but the one rule that is true no matter what the situation is that the easier it is for the H^+ ion to break free, the stronger the acid will be.

The other factor that affects how easily an acid can dissociate is the concentration of the acid. The lower the concentration is for an acid, the higher the percent dissociation will be. This is because the

forward reaction involves the acid donating a proton to a water molecule. In an acid, there is an overabundance of water molecules, so it is very easy for the acid to find a water molecule to donate to. However, the reverse reaction is the hydronium ion donating a proton to the conjugate base. There are a much smaller number of both hydronium and conjugate base ions in a dilute solution, so the reverse reaction is kinetically hindered from happening. This means that more H_3O^+ ions will stay dissociated.

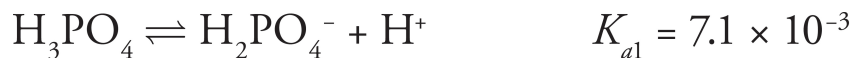
HAt is also considered to be a strong acid, however, given the extremely high reactivity as well as the radioactive nature of astatine, HAt is extremely unstable and of no practical laboratory use.

The greater the concentration of the acid, the more conjugate base there will be, and the easier it will be for that reverse reaction to take place. The easier it is for the reverse reaction to take place, the more HA there will be present in solution, and the less H_3O^+ , leading to a lower overall percent dissociation.

Polyprotic Acids

Some acids, such as H_2SO_4 and H_3PO_4 , can give up more than one hydrogen ion in solution. These are called **polyprotic** acids.

Polyprotic acids are always willing to give up their first proton more easily than future protons. Looking at the three K_a values for phosphoric acid:

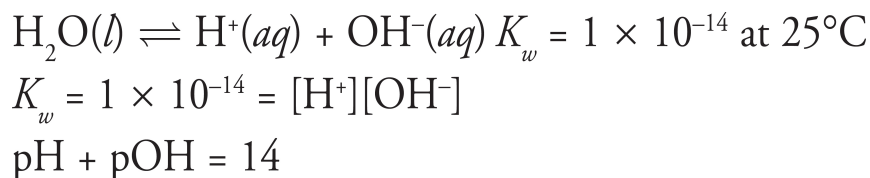


So, H_3PO_4 is a stronger acid than H_2PO_4^- , which in turn is a stronger acid than HPO_4^{2-} . The reason behind this is that after the first proton dissociates, there are a bunch of extra protons floating around in solution. Via Le Châtelier's principle, in future dissociations this causes shifts to the side with no protons—in this case, to the left. That makes it more difficult for remaining protons to dissociate, weakening the acid.

This also means that the amount of each succeeding acid would decrease. In a solution of H_3PO_4 , $[\text{H}_3\text{PO}_4] > [\text{H}_2\text{PO}_4^-] > [\text{HPO}_4^{2-}] > [\text{PO}_4^{3-}]$.

The Equilibrium Constant of Water (K_w)

Water comes to equilibrium with its ions according to the following reaction:



The common ion effect tells us that the hydrogen ion and hydroxide ion concentrations for any acid or base solution must be consistent with the equilibrium for the ionization of water. That is, no matter where the H^+ and OH^- ions came from, when you multiply $[\text{H}^+]$ and $[\text{OH}^-]$, you must get 1×10^{-14} . So for any aqueous solution, if you know the value of $[\text{H}^+]$, you can find out the value of $[\text{OH}^-]$ and vice versa.

The acid and base dissociation constants for conjugates must also be consistent with the equilibrium for the ionization of water.

$$\begin{aligned}K_w &= 1 \times 10^{-14} = K_a K_b \\ \text{p}K_a + \text{p}K_b &= 14\end{aligned}$$

So if you know K_a as a weak acid, you can find K_b for its conjugate base and vice versa.

It is worth noting that pH is not limited to a 0 to 14 range. While most substances will fall into that range, substances that are very acidic can have negative pHs, and substances that are very basic can have pHs that are greater than 14. While you won't find these extreme pHs in most everyday substances, they do exist.

Most acids that you would find in an acids cabinet of a chemistry storeroom are very concentrated and may have negative pH values. For instance, nitric acid is typically manufactured at a concentration of 16 M . Doing the math:

$$-\log(16) = -1.2$$

Needless to say, you should exercise extreme caution when working with concentrated acids or bases, and it should be done only while wearing the proper personal protective equipment.

Something that is important to note is that the K_w for water is 1.0×10^{-14} at a temperature of 25°C only. Like any equilibrium constant, it will change if the temperature does. In this case, the dissociation of water is an endothermic process, as bonds are being broken without any new bonds being formed. So, as temperature increases, the reaction shifts to the right, which increases the value for K_w .

This will have an effect on the pH of water. For instance, at 50°C, $K_w = 5.48 \times 10^{-14}$. Thus, $[H^+] = [OH^-] = 2.34 \times 10^{-7} M$ and $pH = pOH = 6.63$. So, the pH of pure water at 50°C is NOT 7. The K_w of water is often measured by looking at the pK_w value ($-\log K_w$). As temperature and K_w increases, pK_w decreases, as illustrated in the table below.

Temperature (°C)			pH
------------------	--	--	----

	K_w	pK_w	
0	1.14×10^{-15}	14.94	7.47
10	6.81×10^{-15}	14.17	7.27
25	1.00×10^{-14}	14.00	7.00
50	5.48×10^{-14}	13.26	6.63
100	5.13×10^{-13}	12.29	6.14

NEUTRALIZATION REACTIONS

When an acid and a base mix, the acid will donate protons to the base in what is called a neutralization reaction. There are four different mechanisms for this, depending on the strengths of the acids and bases.

1. Strong acid + strong base

When a strong acid mixes with a strong base, both substances are dissociated completely. The only important ions in this type of reaction are the hydrogen and hydroxide ions (Even though not all bases have hydroxides, all strong bases do!).

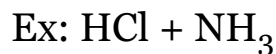
Ex: $\text{HCl} + \text{NaOH}$

Net ionic: $\text{H}^+ + \text{OH}^- \leftrightarrow \text{H}_2\text{O}(l)$

The net ionic equation for all strong acid/strong base reactions is identical—it is always the creation of water. The other ions involved in the reaction (in the example above, Cl^- and Na^+) act as spectator ions and do not take part in the reaction.

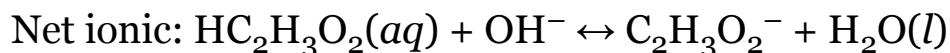
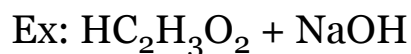
2. Strong acid + weak base

In this reaction, the strong acid (which dissociates completely), will donate a proton to the weak base. The product will be the conjugate acid of the weak base.



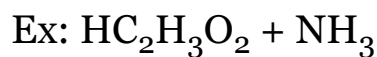
3. Weak acid + strong base

In this reaction, the strong base will accept protons from the weak acid. The products are the conjugate base of the weak acid and water.



4. Weak acid + weak base

This is a simple proton transfer reaction, in which the acid gives protons to the base.



During neutralization reactions, the final pH of the solution is dependent on whether the excess ions at equilibrium are due to the strong acid/base or due to a weak acid/base. When a strong acid/base is in excess, the pH calculation is fairly straightforward.

Example 1: 35.0 mL of 1.5 M HCN, a weak acid ($K_a = 6.2 \times 10^{-10}$) is mixed with 25.0 mL of 2.5 M KOH. Calculate the pH of the final solution.

To do this, we can modify our ICE chart to determine what species are present at equilibrium. First, we must determine the number of moles of the reactants:

$$\text{HCN} = (1.5 \text{ M})(0.035 \text{ L}) = 0.052 \text{ mol} \quad \text{KOH} = (2.5 \text{ M})(0.025 \text{ L}) = 0.062 \text{ mol}$$

Then we set up our ICE chart (leaving values for water out, as it is a pure liquid):

	HCN	OH⁻	CN⁻	H₂O
Initial	0.052	0.062	0	X
Change	-0.052	-0.052	+0.052	X
Equilibrium	0	0.010	0.052	X

The reaction will continue until the HCN runs out of protons to donate to the hydroxide. While there is some weak conjugate base left in solution, it is a weak base and its contribution to the pH of the solution will be irrelevant compared to the strength of the pure hydroxide ions. We now need to determine the concentration of the hydroxide ions at equilibrium. The total volume of the solution is the sum of both the acid and the base. So:

$$\begin{aligned} (35.0 \text{ mL} + 25.0 \text{ mL} &= 60.0 \text{ mL}) \\ [\text{OH}^-] &= 0.010 \text{ mol}/0.060 \text{ L} = 0.17 \text{ M} \\ \text{pOH} &= -\log(0.17) = 0.76 \\ \text{pH} &= 14 - 0.76 = 13.24 \end{aligned}$$

As you can see, it does not take much excess H⁺ or OH⁻ to drive the pH of a solution to a fairly high or low value. Note that the K_a of the acid did not matter in this case, as the strong acid/base was in excess.

When a weak acid or base is in excess, is it easiest to use the Henderson-Hasselbalch equation:

The Henderson-Hasselbalch Equation

$$\text{pH} = \text{p}K_{\text{a}} + \log \frac{[\text{A}^{-}]}{[\text{HA}]}$$

$[\text{HA}]$ = molar concentration of undissociated weak acid (M)

$[\text{A}^{-}]$ = molar concentration of conjugate base (M)

$$\text{pOH} = \text{p}K_{\text{b}} + \log \frac{[\text{HB}^{+}]}{[\text{B}]}$$

$[\text{B}]$ = molar concentration of weak base (M)

$[\text{HB}^{+}]$ = molar concentration of conjugate acid (M)

Example 2: 25.0 mL of 1.0 M HCl is mixed with 60.0 mL of 0.50 M pyridine ($\text{C}_5\text{H}_5\text{N}$), a weak base ($K_{\text{b}} = 1.5 \times 10^{-9}$). Determine the pH of the solution.

Starting out, the number of moles of each reactant must be determined:

$$\text{H}^{+} = (1.0 \text{ } M)(0.025 \text{ L}) = 0.025 \text{ mol H}^{+}$$

$$\text{C}_5\text{H}_5\text{N} = (0.50 \text{ } M)(0.060 \text{ L}) = 0.030 \text{ mol C}_5\text{H}_5\text{N}$$

Then we set up the ICE chart:

--	--	--	--

	H ⁺	C ₅ H ₅ N	HCH ₅ N ⁺
Initial	0.025	0.030	0.000
Change	-0.025	-0.025	+0.025
Equilibrium	0	0.005	0.025

The reaction continues until the H⁺ runs out, leaving pyridine and its conjugate acid at equilibrium. Next, the new concentrations must be determined for both the weak base and its conjugate acid, as both will contribute to the final pH value. The total volume of the new solution is the sum of both the acid and base added, in this case, 25.0 + 60.0 = 85.0 mL. So:

$$[\text{C}_5\text{H}_5\text{N}] = 0.005 \text{ mol} / 0.085 \text{ L} = 0.059 \text{ M}$$

$$[\text{HCH}_5\text{N}^+] = 0.025 \text{ mol} / 0.085 \text{ L} = 0.29 \text{ M}$$

Using Henderson-Hasselbalch:

$$\text{pOH} = \text{p}K_b + \log \frac{[\text{HCH}_5\text{N}^+]}{[\text{C}_5\text{H}_5\text{N}]}$$

$$\text{pOH} = -\log (1.5 \times 10^{-9}) + \log \frac{(0.29)}{(0.059)}$$

$$\text{pOH} = 8.8 + \log (4.9)$$

$$\text{pOH} = 8.8 + 0.69$$

$$\text{pOH} = 9.5$$

$$\text{pH} = 14 - 9.5 = 4.5$$

When a weak acid or base is in excess, the pH of the solution does not change as quickly as when a strong acid or base is in excess.

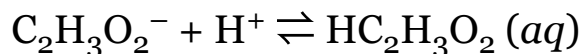
BUFFERS

A **buffer** is a solution with a very stable pH. You can add acid or base to a buffer solution without greatly affecting the pH of the solution. The pH of a buffer will also remain unchanged if the solution is diluted with water or if water is lost through evaporation.

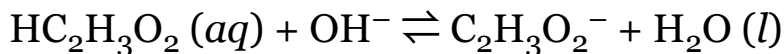
A buffer is created by placing a large amount of a weak acid or base into a solution along with its conjugate, in the form of salt. A weak acid and its conjugate base can remain in solution together without neutralizing each other. This is called the **common ion effect**.

When both the acid and the conjugate base are together in the solution, any hydrogen ions that are added will be neutralized by the base, while any hydroxide ions that are added will be neutralized by the acid, without this having much of an effect on the solution's pH.

If some sodium acetate, $\text{NaC}_2\text{H}_3\text{O}_2$, is added to a solution of acetic acid, $\text{HC}_2\text{H}_3\text{O}_2$, a buffer solution is created, as the acetate ion is the conjugate base of acetic acid. If such a buffer were created and some strong acid were added to it, the protons from the strong acid would react with the acetate ion to create more acetic acid:



Likewise, if a strong base were added to the buffer, the hydroxide ions from the strong base would react with the acetic acid to create more acetate ion:



Through this, we can see that the presence of the conjugate pair is what makes the buffer effective. It's not infinitely effective, of course—if enough strong acid or base is added that ALL of either the acid or its conjugate base is reacted, the buffer will “break.” Thus, the higher the concentration of the conjugate pair, more effectively the more effective the buffer will be. A solution of 1.0 M $\text{HC}_2\text{H}_3\text{O}_2$ and 1.0 M

$\text{NaC}_2\text{H}_3\text{O}_2$ will resist pH change (aka “be a better buffer”) than a solution of $0.10\text{ M HC}_2\text{H}_3\text{O}_2$ and $0.10\text{ M NaC}_2\text{H}_3\text{O}_2$.

From a calculation standpoint, the Henderson-Haselbalch equation allows us to calculate the exact pH of a buffer. If in our acetic acid buffer, $[\text{HC}_2\text{H}_3\text{O}_2] = 0.20\text{ M}$ and $[\text{C}_2\text{H}_3\text{O}_2^-] = 0.50\text{ M}$, the pH of the buffer can be calculated using the below calculations:

$$\begin{aligned}\text{pH} &= \text{p}K_a + \log \frac{[\text{C}_2\text{H}_3\text{O}_2^-]}{[\text{HC}_2\text{H}_3\text{O}_2]} \\ \text{pH} &= -\log (1.8 \times 10^{-5}) + \log \frac{(0.50\text{ M})}{(0.20\text{ M})} \\ \text{pH} &= -\log (1.8 \times 10^{-5}) + \log (2.5) \\ \text{pH} &= (4.7) + (0.40) = 5.1\end{aligned}$$

Now let’s see what happens when $[\text{HC}_2\text{H}_3\text{O}_2]$ and $[\text{C}_2\text{H}_3\text{O}_2^-]$ are both equal to 0.20 M .

$$\begin{aligned}\text{pH} &= \text{p}K_a + \log \frac{[\text{C}_2\text{H}_3\text{O}_2^-]}{[\text{HC}_2\text{H}_3\text{O}_2]} \\ \text{pH} &= -\log (1.8 \times 10^{-5}) + \log \frac{(0.20\text{ M})}{(0.20\text{ M})} \\ \text{pH} &= -\log (1.8 \times 10^{-5}) + \log (1) \\ \text{pH} &= (4.7) + (0) = 4.7\end{aligned}$$

Notice that when the concentrations of acid and conjugate base in a solution are the same, $\text{pH} = \text{p}K_a$ (and $\text{pOH} = \text{p}K_b$). When you choose an acid for a buffer solution, it is best to pick an acid with a $\text{p}K_a$ that is close to the desired pH. That way you can have almost equal amounts of acid and conjugate base in the solution, which will make

the buffer as flexible as possible in neutralizing both added H^+ and OH^- .

That being said, any buffer in which the concentration of the weak acid is greater than the concentration of the conjugate base will logically have a pH lower than the $\text{p}K_a$ of the acid. Any buffer in which the concentration of the conjugate base exceeds the concentration of the weak acid will have a pH that is greater than the $\text{p}K_a$ of the acid. Note that buffers CAN be created from a weak base (such as NH_3) and its conjugate acid (NH_4^+), and although these are far less common the same sort of logic applies as to the weak acid/conjugate base buffers discussed above.

However, you cannot create a buffer solution from a strong acid and its conjugate, because the conjugate base of a strong acid will be very weak. Taking HCl as an example; the Cl^- ion that is left after the acid dissociates completely is a very weak base and will not readily accept protons. The same is true for strong bases; you cannot form a buffer from a strong base and its conjugate for similar reasons.

INDICATORS

Indicators are weak acids which change colors in certain pH ranges due to Le Châtelier's principle. All indicators are weak acids in their own right, but often have very long, very complicated chemical formulas. For simplicity's sake, we will use the generic HIn to indicate a protonated indicator.



$$K_a = \frac{[\text{H}^+][\text{In}^-]}{[\text{HIn}]}$$

For an indicator to be effective, the protonated (HIn) state has to be a different color than the deprotonated (In^-) state. When an indicator is present in a titration and the environment is acidic, the

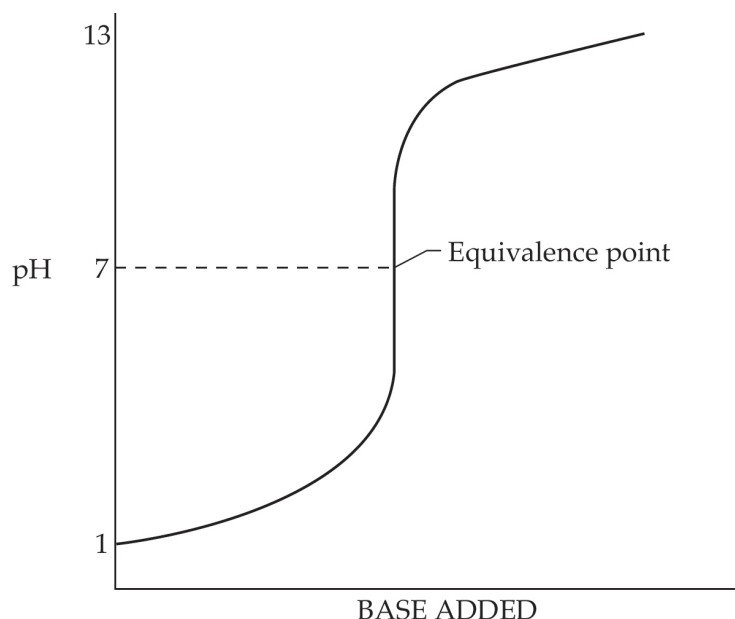
excess H^+ ions drive the equilibrium to the left, causing the solution to consist primarily of HIn and take on that color. In a basic environment, the excess OH^- ions react with the H^+ from the indicator and drive the reaction right, causing the solution to consist of primarily In^- , changing color.

The indicator will experience a color shift right around the point that the two species (HIn and In^-) are present in equal amounts. When that is the case, those two values cancel out of the equilibrium expression, leaving $K_a = [\text{H}^+]$. A quick logarithm later and we can see that at the point of color change, $\text{p}K_a = \text{pH}$. This means that the indicator will change colors at a pH that matches the $\text{p}K_a$ of the indicator.

When doing a titration, you only want to add a few drops of any indicator in order to avoid changing any kind of calculations involving the acid you are titrating against (or with!). It is always best to choose an indicator whose $\text{p}K_a$ matches the pH at the equivalence point of the titration you are doing in order to be able to find the most accurate endpoint.

TITRATION

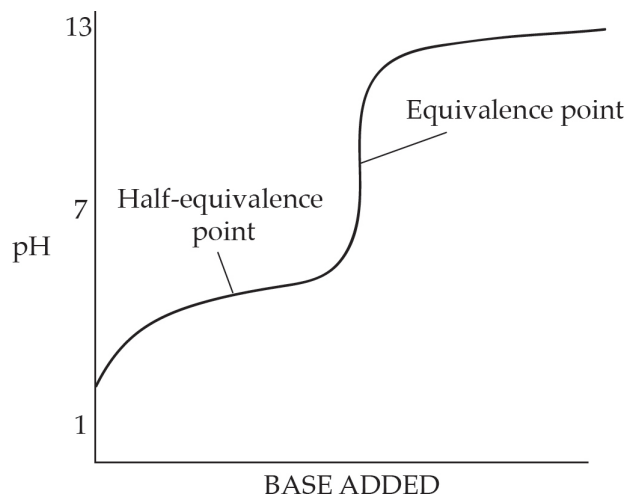
Neutralization reactions are generally performed by titration, where a base of known concentration is slowly added to an acid (or vice versa). The progress of a neutralization reaction can be shown in a titration curve. The diagram below shows the titration of a strong acid by a strong base.



For this titration, the pH at the equivalence point is exactly 7 because the titration of a strong acid by a strong base produces a neutral salt solution.

In the diagram above, the pH increases slowly but steadily from the beginning of the titration until just before the equivalence point. The **equivalence point** is the point in the titration when exactly enough base has been added to neutralize all the acid that was initially present. Just before the equivalence point, the pH increases sharply as the last of the acid is neutralized. The equivalence point of a titration can be recognized through the use of an indicator.

The following diagram shows the titration of a weak acid by a strong base:



In this diagram, the pH increases more quickly at first, then levels out into a buffer region. At the center of the buffer region is the **half-equivalence point**.

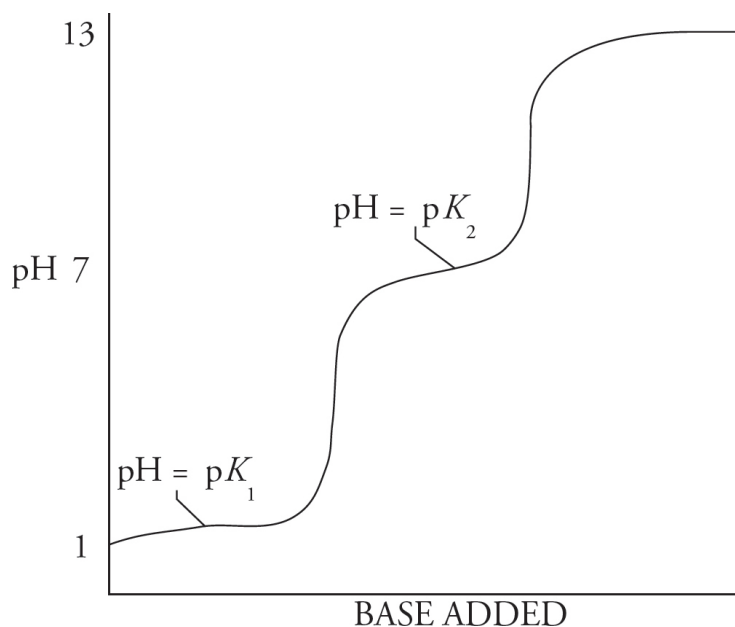
At this point, enough base has been added to convert exactly half of the acid into conjugate base; here the concentration of acid is equal to the concentration of conjugate base $[HA] = [A^-]$. Putting that into our weak acid dissociation expression:

$$K_a = \frac{[H^+][A^-]}{[HA]}$$

If $[A^-]$ and $[HA]$ are equal and cancel out, that leaves us with $K_a = [H^+]$, which is often represented by $pK_a = pH$. This is a good way to determine the K_a of a weak acid from a titration curve such as the one above.

The curve remains fairly flat until just before the equivalence point, when the pH increases sharply. For this titration, the pH at the equivalence point is greater than 7 because the only ion present in significant amounts at equilibrium is the conjugate base of the weak acid. Likewise, if you titrate a weak base with a strong acid, the solution will be acidic at equilibrium because the only ion present in significant amounts will be the conjugate acid of the weak base.

The following diagram shows the titration curve of a polyprotic acid:



For a polyprotic acid, the titration curve will have as many bumps as there are hydrogen ions to give up. The curve above has two bumps, so it represents the titration of a diprotic acid. There are also two half-equivalence points for a diprotic acid. At the first one, the pH will be equal to the negative log of the first dissociation constant for the polyprotic acid (pK_1). At the second half-equivalence point, the pH will be equal to the negative log of the second dissociation constant (pK_2).

CHAPTER 8 QUESTIONS

[Click here](#) to download and print the PDF version of this exercise.

Multiple-Choice

Use the following information to answer questions 1-5.

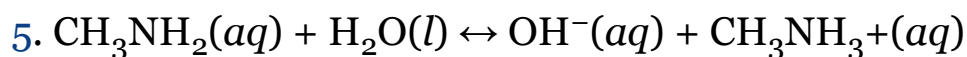
A student titrates 20.0 mL of 1.0 *M* NaOH with 2.0 *M* formic acid, HCO_2H ($K_a = 1.8 \times 10^{-4}$). Formic acid is a monoprotic acid.

1. How much formic acid is necessary to reach the equivalence point?
 - (A) 10.0 mL
 - (B) 20.0 mL
 - (C) 30.0 mL
 - (D) 40.0 mL
2. At the equivalence point, is the solution acidic, basic, or neutral? Why?
 - (A) Acidic; the strong acid dissociates more than the weak base
 - (B) Basic; the only ion present at equilibrium is the conjugate base
 - (C) Basic; the higher concentration of the base is the determining factor
 - (D) Neutral; equal moles of both acid and base are present
3. If the formic acid were replaced with a strong acid such as HCl at the same concentration (2.0 *M*), how would that change the

volume needed to reach the equivalence point?

- (A) The change would reduce the amount as the acid now fully dissociates.
 - (B) The change would reduce the amount because the base will be more strongly attracted to the acid.
 - (C) The change would increase the amount because the reaction will now go to completion instead of equilibrium.
 - (D) Changing the strength of the acid will not change the volume needed to reach equivalence.
4. Which of the following would create a good buffer when dissolved in formic acid?

- (A) NaCO_2H
- (B) $\text{HC}_2\text{H}_3\text{O}_2$
- (C) NH_3
- (D) H_2O



The above equation represents the reaction between the base methylamine ($K_b = 4.38 \times 10^{-4}$) and water. Which of the following best represents the concentrations of the various species at equilibrium?

- (A) $[\text{OH}^-] > [\text{CH}_3\text{NH}_2] = [\text{CH}_3\text{NH}_3^+]$
- (B) $[\text{OH}^-] = [\text{CH}_3\text{NH}_2] = [\text{CH}_3\text{NH}_3^+]$
- (C) $[\text{CH}_3\text{NH}_2] > [\text{OH}^-] > [\text{CH}_3\text{NH}_3^+]$
- (D) $[\text{CH}_3\text{NH}_2] > [\text{OH}^-] = [\text{CH}_3\text{NH}_3^+]$

Use the following information to answer questions 6-10.

The following reaction is found to be at equilibrium at 25°C:



6. What is the expression for the equilibrium constant, K_c ?

- (A) $\frac{[\text{SO}_3]^2}{[\text{O}_2][\text{SO}_2]^2}$
- (B) $\frac{2[\text{SO}_3]}{[\text{O}_2]2[\text{SO}_2]}$
- (C) $\frac{[\text{O}_2][\text{SO}_2]^2}{[\text{SO}_3]^2}$
- (D) $\frac{[\text{O}_2]2[\text{SO}_2]}{2[\text{SO}_3]}$

7. Which of the following would cause the reverse reaction to speed up?

- (A) Adding more SO_3
- (B) Raising the pressure
- (C) Lowering the temperature
- (D) Removing some SO_2

8. The value for K_c at 25°C is 8.1. What must happen in order for the reaction to reach equilibrium if the initial concentrations of all three species was 2.0 M?

- (A) The rate of the forward reaction would increase, and $[\text{SO}_3]$ would decrease.

- (B) The rate of the reverse reaction would increase, and $[\text{SO}_2]$ would decrease.
- (C) Both the rate of the forward and reverse reactions would increase, and the value for the equilibrium constant would also increase.
- (D) No change would occur in either the rate of reaction or the concentrations of any of the species.
9. Which of the following would cause a reduction in the value for the equilibrium constant?
- (A) Increasing the amount of SO_3
- (B) Reducing the amount of O_2
- (C) Raising the temperature
- (D) Lowering the temperature
10. The solubility product, K_{sp} , of AgCl is 1.8×10^{-10} . Which of the following expressions is equal to the solubility of AgCl ?
- (A) $(1.8 \times 10^{-10})^2$ molar
- (B) $\frac{1.8 \times 10^{-10}}{2}$ molar
- (C) 1.8×10^{-10} molar
- (D) $\sqrt{1.8 \times 10^{-10}}$ molar
11. A 0.1-molar solution of which of the following acids will be the best conductor of electricity?
- (A) H_2CO_3
- (B) H_2S
- (C) HF

(D) HNO_3

12. Which of the following expressions is equal to the K_{sp} of Ag_2CO_3 ?

(A) $K_{sp} = [\text{Ag}^+][\text{CO}_3^{2-}]$

(B) $K_{sp} = [\text{Ag}^+][\text{CO}_3^{2-}]^2$

(C) $K_{sp} = [\text{Ag}^+]^2[\text{CO}_3^{2-}]$

(D) $K_{sp} = [\text{Ag}^+]^2[\text{CO}_3^{2-}]^2$

13. If the solubility of BaF_2 is equal to x , which of the following expressions is equal to the solubility product, K_{sp} , for BaF_2 ?

(A) x^2

(B) $2x^2$

(C) $2x^3$

(D) $4x^3$

Use the following information to answer questions 14-16:

150 mL of saturated SrF_2 solution is present in a 250 mL beaker at room temperature. The molar solubility of SrF_2 at 298 K is $1.0 \times 10^{-3} \text{ M}$.

14. What are the concentrations of Sr^{2+} and F^- in the beaker?

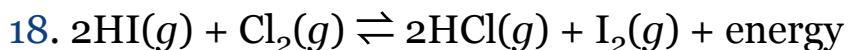
(A) $[\text{Sr}^{2+}] = 1.0 \times 10^{-3} \text{ M}$ $[\text{F}^-] = 1.0 \times 10^{-3} \text{ M}$

(B) $[\text{Sr}^{2+}] = 1.0 \times 10^{-3} \text{ M}$ $[\text{F}^-] = 2.0 \times 10^{-3} \text{ M}$

(C) $[\text{Sr}^{2+}] = 2.0 \times 10^{-3} \text{ M}$ $[\text{F}^-] = 1.0 \times 10^{-3} \text{ M}$

(D) $[\text{Sr}^{2+}] = 2.0 \times 10^{-3} \text{ M}$ $[\text{F}^-] = 2.0 \times 10^{-3} \text{ M}$

15. If some of the solution evaporates overnight, which of the following will occur?
- (A) The mass of the solid and the concentration of the ions will stay the same.
 - (B) The mass of the solid and the concentration of the ions will increase.
 - (C) The mass of the solid will decrease, and the concentration of the ions will stay the same.
 - (D) The mass of the solid will increase, and the concentration of the ions will stay the same.
16. How could the concentration of Sr^{2+} ions in solution be decreased?
- (A) Adding some NaF(s) to the beaker
 - (B) Adding some $\text{Sr(NO}_3)_2\text{(s)}$ to the beaker
 - (C) By heating the solution in the beaker
 - (D) By adding a small amount of water to the beaker, but not dissolving all the solid
17. For a reaction involving nitrogen monoxide inside a sealed flask, the value for the reaction quotient (Q) was found to be 1.1×10^2 at a given point. If, after this point, the amount of NO gas in the flask increased, which reaction is most likely taking place in the flask?
- (A) $\text{NOBr(g)} \leftrightarrow \text{NO(g)} + \frac{1}{2}\text{Br}_2\text{(g)}$ $K_c = 3.4 \times 10^{-2}$
 - (B) $2\text{NOCl(g)} \leftrightarrow 2\text{NO(g)} + \text{Cl}_2\text{(g)}$ $K_c = 1.6 \times 10^{-5}$
 - (C) $2\text{NO(g)} + 2\text{H}_2\text{(g)} \leftrightarrow \text{N}_2\text{(g)} + 2\text{H}_2\text{O(g)}$ $K_c = 4.0 \times 10^6$
 - (D) $\text{N}_2\text{(g)} + \text{O}_2\text{(g)} \leftrightarrow 2\text{NO(g)}$ $K_c = 4.2 \times 10^2$



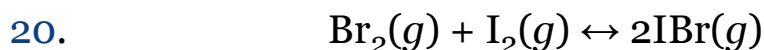
A gaseous reaction occurs and comes to equilibrium, as shown above. Which of the following changes to the system will serve to increase the number of moles of I_2 present at equilibrium?

- (A) Increasing the volume at constant temperature
- (B) Decreasing the volume at constant temperature
- (C) Increasing the temperature at constant volume
- (D) Decreasing the temperature at constant volume

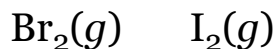


The reaction above came to equilibrium at a temperature of 100°C . At equilibrium the partial pressure due to NOBr was 4 atmospheres, the partial pressure due to NO was 4 atmospheres, and the partial pressure due to Br_2 was 2 atmospheres. What is the equilibrium constant, K_p , for this reaction at 100°C ?

- (A) $\frac{1}{4}$
- (B) $\frac{1}{2}$
- (C) 1
- (D) 2



At 150°C, the equilibrium constant, K_c , for the reaction shown above has a value of 300. This reaction was allowed to reach equilibrium in a sealed container and the partial pressure due to $\text{IBr}(g)$ was found to be 3 atm. Which of the following could be the partial pressures due to $\text{Br}_2(g)$ and $\text{I}_2(g)$ in the container?



- (A) 0.1 atm 0.3 atm
- (B) 0.3 atm 1 atm
- (C) 1 atm 1 atm
- (D) 1 atm 3 atm

21. A laboratory technician wishes to create a buffered solution with a pH of 5. Which of the following acids would be the best choice for the buffer?

- | | |
|--|----------------------------|
| (A) $\text{H}_2\text{C}_2\text{O}_4$ | $K_a = 5.9 \times 10^{-2}$ |
| (B) H_3AsO_4 | $K_a = 5.6 \times 10^{-3}$ |
| (C) $\text{H}_2\text{C}_2\text{H}_3\text{O}_2$ | $K_a = 1.8 \times 10^{-5}$ |
| (D) HOCl | $K_a = 3.0 \times 10^{-8}$ |

22. Which of the following species is amphoteric?

- (A) H^+
- (B) CO_3^{2-}
- (C) HCO_3^-
- (D) H_2CO_3

23. How many liters of distilled water must be added to 1 liter of an aqueous solution of HCl with a pH of 1 to create a solution with a pH of 2?

- (A) 0.1 L
- (B) 0.9 L
- (C) 2 L
- (D) 9 L

24. A 1-molar solution of a very weak monoprotic acid has a pH of 5. What is the value of K_a for the acid?

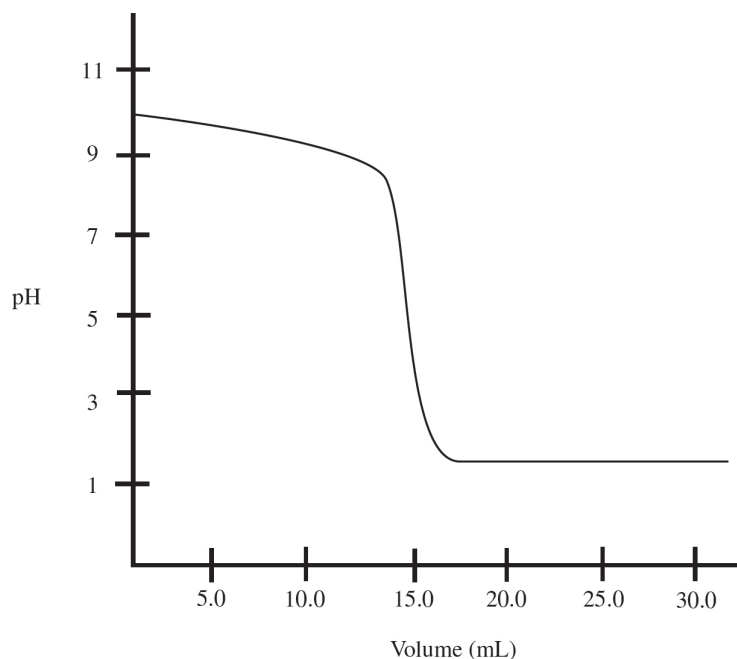
- (A) $K_a = 1 \times 10^{-10}$
- (B) $K_a = 1 \times 10^{-7}$
- (C) $K_a = 1 \times 10^{-5}$
- (D) $K_a = 1 \times 10^{-2}$

25. The value of K_a for HSO_4^- is 1×10^{-2} . What is the value of K_b for SO_4^{2-} ?

- (A) $K_b = 1 \times 10^{-12}$
- (B) $K_b = 1 \times 10^{-8}$
- (C) $K_b = 1 \times 10^{-2}$
- (D) $K_b = 1 \times 10^2$

Use the following information to answer questions 26-28.

The following curve is obtained during the titration of 30.0 mL of 1.0 M NH_3 , a weak base, with a strong acid:



26. Why is the solution acidic at equilibrium?

- (A) The strong acid dissociates fully, leaving excess $[\text{H}^+]$ in solution.
- (B) The conjugate acid of NH_3 is the only ion present at equilibrium.
- (C) The water which is being created during the titration acts as an acid.
- (D) The acid is diprotic, donating two protons for every unit dissociated.

27. What is the concentration of the acid?

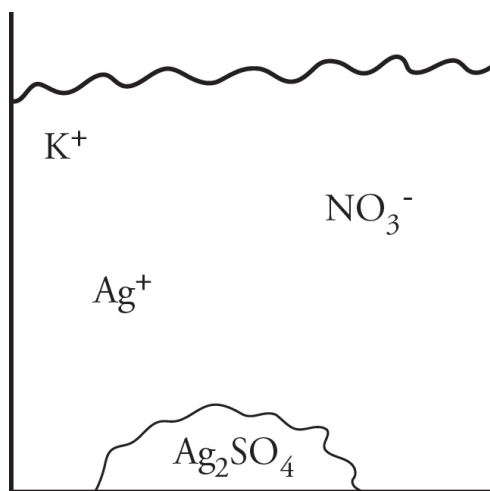
- (A) 0.5 *M*
- (B) 1.0 *M*
- (C) 1.5 *M*
- (D) 2.0 *M*

28. What ions are present in significant amounts during the first buffer region?

- (A) NH_3 and NH_4^+
- (B) NH_3 and H^+
- (C) NH_4^+ and OH^-
- (D) H_3O^+ and NH_3

Use the information below to answer questions 29-31.

Silver sulfate, Ag_2SO_4 , has a solubility product constant of 1.0×10^{-5} . The below diagram shows the products of a precipitation reaction in which some silver sulfate was formed.



29. What is the identity of the excess reactant?

- (A) AgNO_3
- (B) Ag_2SO_4
- (C) NaNO_3
- (D) Na_2SO_4

30. If the beaker above were left uncovered for several hours:
- (A) Some of the Ag_2SO_4 would dissolve.
 - (B) Some of the spectator ions would evaporate into the atmosphere.
 - (C) The solution would become electrically imbalanced.
 - (D) Additional Ag_2SO_4 would precipitate.
31. Which ion concentrations below would have led the precipitate to form?
- (A) $[\text{Ag}^+] = 0.01\text{ M}$ $[\text{SO}_4^{2-}] = 0.01\text{ M}$
 - (B) $[\text{Ag}^+] = 0.10\text{ M}$ $[\text{SO}_4^{2-}] = 0.01\text{ M}$
 - (C) $[\text{Ag}^+] = 0.01\text{ M}$ $[\text{SO}_4^{2-}] = 0.10\text{ M}$
 - (D) This is impossible to determine without knowing the total volume of the solution.
32. In a voltaic cell with a $\text{Cu (s)} \mid \text{Cu}^{2+}$ cathode and a $\text{Pb}^{2+} \mid \text{Pb (s)}$ anode, increasing the concentration of Pb^{2+} causes the voltage to decrease. What is the reason for this?
- (A) The value for Q will increase, causing the cell to come closer to equilibrium.
 - (B) The solution at the anode becomes more positively charged, leading to a reduced electron flow.
 - (C) The reaction will shift to the right, causing a decrease in favorability.
 - (D) Cell potential will always decrease anytime the concentration of any aqueous species present increases.
33. Which of the following could be added to an aqueous solution of weak acid HF to increase the percent dissociation?

- (A) NaF(s)
- (B) $\text{H}_2\text{O(l)}$
- (C) NaOH(s)
- (D) $\text{NH}_3(\text{aq})$

34. A bottle of water is left outside early in the morning. The bottle warms gradually over the course of the day. What will happen to the pH of the water as the bottle warms?

- (A) Nothing; pure water always has a pH of 7.00.
- (B) Nothing; the volume would have to change in order for any ion concentration to change.
- (C) It will increase because the concentration of $[\text{H}^+]$ is increasing.
- (D) It will decrease because the auto-ionization of water is an endothermic process.

35. In which of the following circumstances is the value for K_{eq} always greater than 1?

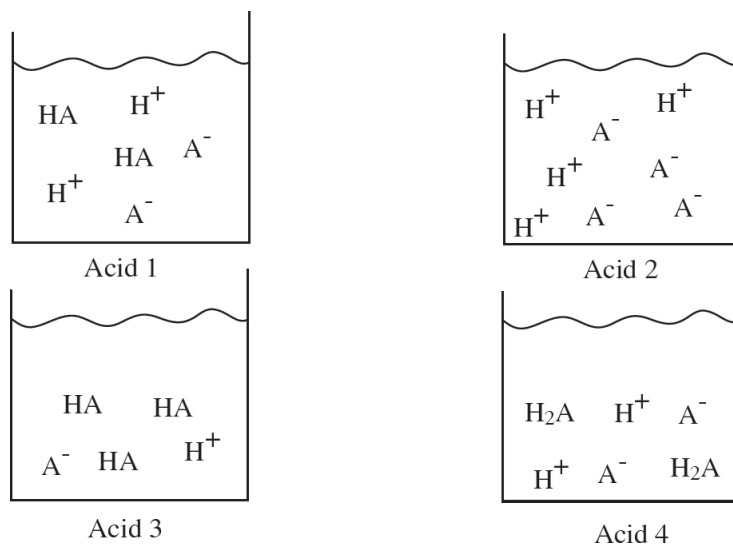
	ΔH	ΔS
(A)	Positive	Positive
(B)	Positive	Negative
(C)	Negative	Negative
(D)	Negative	Positive

36. The structure of two oxoacids is shown below:



Which would be a stronger acid, and why?

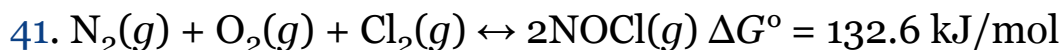
- (A) HOCl, because the H–O bond is weaker than in HOF as chlorine is larger than fluorine
- (B) HOCl, because the H–O bond is stronger than in HOF as chlorine has a higher electronegativity than fluorine
- (C) HOF, because the H–O bond is stronger than in HOCl as fluorine has a higher electronegativity than chlorine
- (D) HOF, because the H–O bond is weaker than in HOCl as fluorine is smaller than chlorine
37. Which of the following pairs of substances would make a good buffer solution?
- (A) $\text{HC}_2\text{H}_3\text{O}_2(aq)$ and $\text{NaC}_2\text{H}_3\text{O}_2(aq)$
- (B) $\text{H}_2\text{SO}_4(aq)$ and $\text{LiOH}(aq)$
- (C) $\text{HCl}(aq)$ and $\text{KCl}(aq)$
- (D) $\text{HF}(aq)$ and $\text{NH}_3(aq)$
38. $\text{PCl}_3(g) + \text{Cl}_2(g) \leftrightarrow \text{PCl}_5(g) \quad \Delta H = -92.5 \text{ kJ/mol}$
- In which of the following ways could the reaction above be manipulated to create more product?
- (A) Decreasing the concentration of PCl_3
- (B) Increasing the pressure
- (C) Increasing the temperature
- (D) None of the above
39. Four different acids are added to beakers of water, and the following diagrams represent the species present in each solution at equilibrium. Which acid has the highest pH?



- (A) Acid 1
- (B) Acid 2
- (C) Acid 3
- (D) Acid 4

40. Starting with a stock solution of $18.0\text{ M H}_2\text{SO}_4$, what is the proper procedure to create a 1.00 L sample of a 3.0 M solution of H_2SO_4 in a volumetric flask?

- (A) Add 167 mL of the stock solution to the flask, then fill the flask the rest of the way with distilled water while swirling the solution.
- (B) Add 600 mL of the stock solution to the flask, then fill the flask the rest of the way with distilled water while swirling the solution.
- (C) Fill the flask partway with water, then add 167 mL of the stock solution, swirling to mix it. Last, fill the flask the rest of the way with distilled water.
- (D) Fill the flask partway with water, then add 600 mL of the stock solution, swirling to mix it. Last, fill the flask the rest of the way with distilled water.



For the equilibrium above, what would happen to the value of ΔG° if the concentration of N_2 were to increase and why?

- (A) It would increase because the reaction would become more thermodynamically favored.
- (B) It would increase because the reaction would shift right and create more products.
- (C) It would decrease because there are more reactants present.
- (D) It would stay the same because the value of K_{eq} would not change.

Free-Response Questions

1. The value of the solubility product, K_{sp} , for calcium hydroxide, $\text{Ca}(\text{OH})_2$, is 5.5×10^{-6} , at 25°C .
 - (a) Write the K_{sp} expression for calcium hydroxide.
 - (b) What is the mass of $\text{Ca}(\text{OH})_2$ in 500 mL of a saturated solution at 25°C ?
 - (c) What is the pH of the solution in (b)?
 - (d) If 1.0 mole of OH^- is added to the solution in (b), what will be the resulting Ca^{2+} concentration? Assume that the volume of the solution does not change.
2. For sodium chloride, the solution process with water is endothermic.
 - (a) Describe the change in entropy when sodium chloride dissociates into aqueous particles.

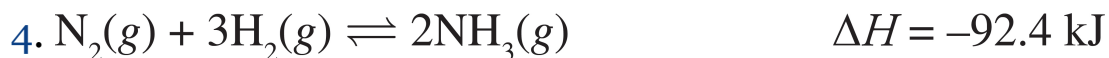
- (b) Two saturated aqueous NaCl solutions, one at 20°C and one at 50°C, are compared. Which one will have higher concentration? Justify your answer.
- (c) Which way will the solubility reaction shift if the temperature is increased?
- (d) If a saturated solution of NaCl is left out overnight and some of the solution evaporates, how will that affect the amount of solid NaCl present?
3. A student tests the conductivity of three different acid samples, each with a concentration of 0.10 *M* and a volume of 20.0 mL. The conductivity was recorded in microsiemens per centimeter in the table below:

Sample	Conductivity ($\mu\text{S}/\text{cm}$)
1	26,820
2	8655
3	35,120

- (a) The three acids are known to be HCl, H₂SO₄, and H₃PO₄. Identify which sample is which acid. Justify your answer.
- (b) The HCl solution is then titrated with a 0.150 *M* solution of the weak base methylamine, CH₃NH₂.

$$(K_b = 4.38 \times 10^{-4})$$

- (i) Write out the net ionic equation for this reaction.
- (ii) Determine the pH of the solution after 20.0 mL of methylamine has been added.



When the reaction above took place at a temperature of 570 K, the following equilibrium concentrations were measured:

$$[\text{NH}_3] = 0.20 \text{ mol/L}$$

$$[\text{N}_2] = 0.50 \text{ mol/L}$$

$$[\text{H}_2] = 0.20 \text{ mol/L}$$

- (a) Write the expression for K_c and calculate its value.
- (b) Calculate ΔG for this reaction.
- (c) Describe how the concentration of H_2 at equilibrium will be affected by each of the following changes to the system at equilibrium:
 - (i) The temperature is increased.
 - (ii) The volume of the reaction chamber is increased.
 - (iii) N_2 gas is added to the reaction chamber.
 - (iv) Helium gas is added to the reaction chamber.

5. In an acidic medium, iron (III) ions will react with thiocyanate (SCN^-) ions to create the following complex ion:



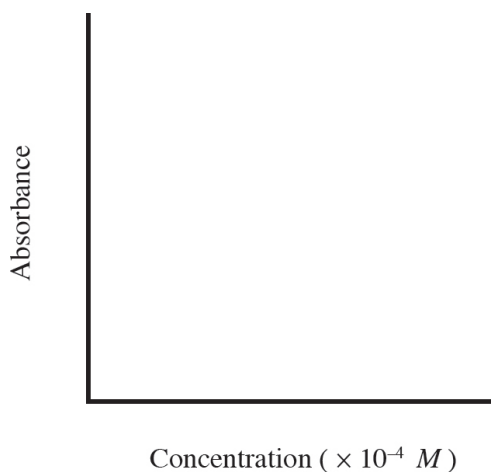
Initially, the solution is a light yellow color due to the presence of the Fe^{3+} ions. As the FeSCN^{2+} forms, the solution will gradually darken to a golden yellow. The reaction is not a fast one, and generally after mixing the ions the maximum concentration of FeSCN^{2+} will occur between 2–4 minutes after mixing the solution.

A student creates four solutions with varying concentration of FeSCN^{2+} and gathers the following data at 298 K using a

spectrophotometer calibrated to 460 nm:

[FeSCN ²⁺]	Absorbance
$1.1 \times 10^{-4} M$	0.076
$1.6 \times 10^{-4} M$	0.112
$2.2 \times 10^{-4} M$	0.167
$2.5 \times 10^{-4} M$	0.199

- (a) (i) On the axes below, create a Beer's Law calibration plot for [FeSCN²⁺]. Draw a best-fit line through your data points.



- (ii) The slope of the best-fit line for the above set of data points is 879 and the y -intercept is -0.024 . Write out the equation for this line.

To determine the equilibrium constant for the reaction, a solution is made up in which 5.00 mL of 0.0025 M Fe(NO₃)₃ and 5.00 mL of 0.0025 M KSCN are mixed. After 3 minutes, the absorbance of the solution is found to be 0.134.

- (b) (i) Using your Beer's Law best-fit line from (a), calculate [FeSCN²⁺] once equilibrium has been established.

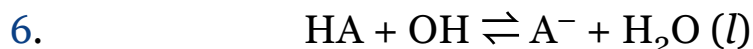
(ii) Calculate $[\text{Fe}^{3+}]$ and $[\text{SCN}^-]$ at equilibrium.

(iii) Calculate K_{eq} for the reaction.

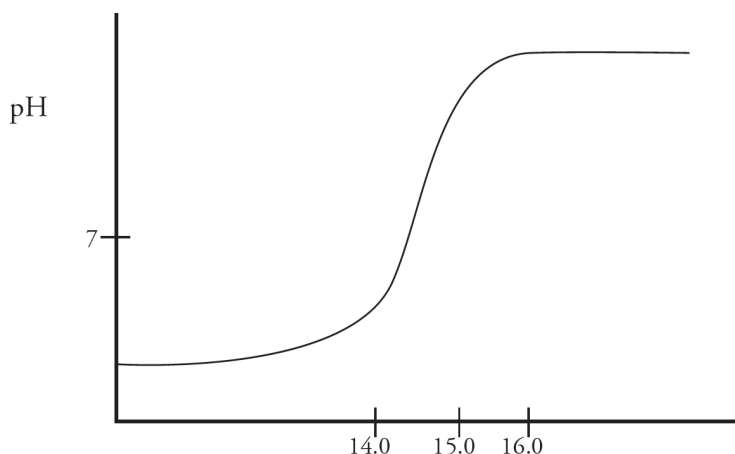
After equilibrium is established, the student heats the solution and observes that it becomes noticeably lighter.

(c) (i) Did heating the mixture increase the equilibrium constant, decrease it, or have no effect on it? Why?

(ii) Is the equilibrium reaction exothermic or endothermic? Justify your answer.



A student titrates a weak acid, HA, with some 1.0 M NaOH, yielding the following titration curve:



(a) Which chemical species present in solution dictates the pH of the solution in each of the volume ranges listed below?

(i) 1.0 mL–14.0 mL

(ii) 15.0 mL

(iii) 16.0 mL–30.0 mL

(b) At which volumes is:

(i) $[\text{HA}] > [\text{A}^-]$?

(ii) $[HA] = [A^-]$?

(iii) $[HA] < [A^-]$?

(c) At which point in the titration (if any) would the concentration of the following species be equal to zero? Justify your answers.

(i) HA

(ii) A^-

(d) If the titration were performed again, but this time with 2.0 M NaOH, name two things that would change about the titration curve, and explain the reasoning behind your identified changes.

7. A student performs an experiment to determine the concentration of a solution of hypochlorous acid, HOCl ($K_a = 3.5 \times 10^{-8}$). The student starts with 25.00 mL of the acid in a flask and titrates it against a standardized solution of sodium hydroxide with a concentration of 1.47 M. The equivalence point is reached after the addition of 34.23 mL of NaOH.

(a) Write the net ionic equation for the reaction that occurs in the flask.

(b) What is the concentration of the HOCl?

(c) What would the pH of the solution in the flask be after the addition of 28.55 mL of NaOH?

(d) The actual concentration of the HOCl is found to be 2.25 M. Quantitatively discuss whether or not each of the following errors could have caused the error in the student's results.

(i) The student added additional NaOH past the equivalence point.

(ii) The student rinsed the buret with distilled water but not with the NaOH solution before filling it with NaOH.

(iii) The student measured the volume of acid incorrectly; instead of adding 25.00 mL of HOCl, only 24.00 mL was present in the flask prior to titration.

8. A student is tasked with determining the identity of an unknown carbonate compound with a mass of 1.89 g. The compound is first placed in water, where it dissolves completely. The K_{sp} value for several carbonate-containing compounds are given below.

Compound	K_{sp}
Lithium carbonate	8.15×10^{-4}
Nickel (II) carbonate	1.42×10^{-7}
Strontium carbonate	5.60×10^{-10}

- (a) In order to precipitate the maximum amount of the carbonate ions from solution, which of the following should be added to the carbonate solution: LiNO_3 , $\text{Ni(NO}_3)_2$, or $\text{Sr(NO}_3)_2$? Justify your answer.
- (b) For the carbonate compound that contains the cation chosen in part (a), determine the concentration of each ion of that compound in solution at equilibrium.
- (c) When mixing the solution, should the student ensure the carbonate solution or the nitrate solution is in excess? Justify your answer.
- (d) After titrating sufficient solution to precipitate out all of the carbonate ions, the student filters the solution before placing it in a crucible and heating it to drive off the water. After several heatings, the final mass of the precipitate remains constant and is determined to be 2.02 g.
- (i) Determine the number of moles of precipitate.

- (ii) Determine the mass of carbonate present in the precipitate.
- (e) Determine the percent, by mass, of carbonate in the original sample.
- (f) Is the original compound most likely lithium carbonate, sodium carbonate, or potassium carbonate? Justify your answer.

CHAPTER 8 ANSWERS AND EXPLANATIONS

Multiple-Choice

1. **A**

The equivalence point is defined as the point at which the moles of acid are equal to the moles of base. The amount of NaOH is equal to $(1.0\text{ M})(0.0200\text{ L}) = 0.0200\text{ mol}$. To calculate the volume of acid:

$$2.0\text{ M} = 0.0200\text{ mol}/V$$

$$V = 0.0100\text{ L} = 10.0\text{ mL}$$

2. **B**

Using the ICE chart:

	HCO_2H	OH^-	CO_2H^-	H_2O
Initial	0.020	0.020	0	X
Change	-0.020	-0.020	+ 0.020	X
Equilibrium	0	0	0.020	X

Both the hydroxide ion and the weak acid are not present at equilibrium. The only ionic species left, the O_2CH^- , is basic, and so the solution will be as well.

3. **D**

The number of moles of base is staying the same, and so is the concentration of the acid. Therefore, the same volume of that acid will be needed to get to the equivalence point.

4. **A**

A buffer is made up of an acid and its conjugate base. The conjugate base of HCO_2H is CO_2H^- , which is present as the anion in the NaCO_2H salt.

5. **D**

Weak bases do not ionize fully in solution, and most of the methylamine molecules will not deprotonate. The hydroxide and conjugate acid ions are created in a 1:1 ratio and therefore will be equal.

6. **C**

The equilibrium expression is always products over reactants, so that eliminates (A) and (B). The coefficients in the balanced equation turn into exponents in the expression, leading you to the correct answer.

7. **B**

To speed up the reverse reaction, you are looking to cause a shift to the left via Le Châtelier's principle. If the pressure were to increase, there would be a shift to the side with fewer gas molecules, which in this case means a shift to the left. All other options cause a shift to the right.

8. **A**

To determine which way the reaction would shift, the reaction quotient, Q , would need to be calculated. This can be done by plugging the concentrations into the Law of Mass Action. With all of the initial values being 2.0 M :

$$Q = \frac{(2.0)(2.0)^2}{(2.0)^2} = 2.0$$

2.0 < 8.1, so the reaction must shift right in order for the reaction to proceed to equilibrium. This would cause an increase in the rate of the forward reaction, along with a decrease in the $[\text{SO}_3]$.

9. **C**

Changing the amounts of either reactants or products present will not cause a change in the equilibrium constant (nor would changing the pressure, if that were an option). The only way to actually change the value of the constant is by changing the temperature. As this is an exothermic reaction, adding more heat would cause a shift to the left, increasing the amount of reactants, and thus the denominator in the equilibrium constant expression. This, in turn, reduces the value for K_c .

10. **D**

The solubility of a substance is equal to its maximum concentration in solution.

For every AgCl in solution, you get one Ag^+ and one Cl^- , so the solubility of AgCl—let's call it x —will be the same as $[\text{Ag}^+]$, which is the same as $[\text{Cl}^-]$.

So for AgCl, $K_{sp} = [\text{Ag}^+][\text{Cl}^-] = 1.8 \times 10^{-10} = x^2$.

$$x = \sqrt{1.8 \times 10^{-10}}$$

11. **D**

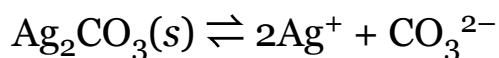
The best conductor of electricity (also called the strongest electrolyte) will be the solution that contains the most charged particles. HNO_3 is the only strong acid listed in the

answer choices, so it is the only choice where the acid has dissociated completely in solution into H^+ and NO_3^- ions. So a 0.1-molar HNO_3 solution will contain the most charged particles and therefore be the best conductor of electricity.

12. **C**

K_{sp} is just the equilibrium constant without a denominator.

When Ag_2CO_3 dissociates, you get the following reaction:



In the equilibrium expression, coefficients become exponents, so you get:

$$K_{sp} = [\text{Ag}^+]^2[\text{CO}_3^{2-}]$$

13. **D**

For BaF_2 , $K_{sp} = [\text{Ba}^{2+}][\text{F}^-]^2$.

For every BaF_2 that dissolves, you get one Ba^{2+} and two F^- .

So if the solubility of BaF_2 is x , then $[\text{Ba}^{2+}] = x$, and $[\text{F}^-] = 2x$.

$$\text{So } K_{sp} = (x)(2x)^2 = (x)(4x^2) = 4x^3$$

14. **B**

When SrF_2 dissociates, it creates one Sr^{2+} ion and two F^- ions. That means the concentration of fluoride ions will be twice that of strontium ions in a saturated solution.

15. **D**

If the solution is saturated, that means the concentrations of Sr^{2+} and F^- are at their maximum values and would remain unchanged even if some water evaporated. If that happens, thus decreasing the volume of the solution, there will not be as much room for the solute ions in the solution. These ions would fall out of solution, increasing the mass of the solid, and maintaining the same concentration of ions in solution.

16. **A**

If extra F^- ions are added to the solution via the addition of NaF (which contains an alkali metal cation and is thus fully soluble), they will bond with some of the Sr^{2+} ions that were present to create more $\text{SrF}_2(\text{s})$ and reduce the number of Sr^{2+} ions present in solution. This is called the common ion effect. Choice (D) is incorrect because as long as there is still solid on the bottom of the beaker, the equilibrium concentration of the ions will remain unchanged.

17. **D**

When $Q > K_c$, the numerator of the equilibrium expression (the product concentration) is too big, and the equation shifts to the left. This is true for both (A) and (B), meaning $[\text{NO}]$ would decrease. When $Q < K_c$, the numerator/product concentrations need to increase. This is the case in (C) and (D), but $\text{NO}(\text{g})$ is only a product in (D).

18. **D**

According to Le Châtelier's Law, the equilibrium will shift to counteract any stress that is placed on it. If the temperature is decreased, the equilibrium will shift toward

the side that produces energy or heat. That's the product side where I_2 is produced.

19. **D**

$$K_p = \frac{[NO]^2 [Br_2]}{[NOBr]^2} = \frac{(4)^2 (2)}{(4)^2} = 2$$

20. **D**

The equilibrium expression for the reaction is as follows:

$$\frac{P_{I_{Br}}^2}{P_{Br_2} P_{I_2}} = 300$$

When all of the values are plugged into the expression, (A) is the only choice that works.

$$\frac{(3)^2}{(0.1)(0.3)} = \frac{9}{0.03} = 300$$

21. **C**

The best buffered solution occurs when $pH = pK_a$. That happens when the solution contains equal amounts of acid and conjugate base. If you want to create a buffer with a pH of 5, the best choice would be an acid with a pK_a that is as close to 5 as possible. You shouldn't have to do a calculation to see that the pK_a for (C) is much closer to 5 than that of any of the others.

22. **C**

An amphoteric species can act either as an acid or a base, gaining or losing a proton.

HCO_3^- can act as an acid, losing a proton to become CO_3^{2-} , or it can act as a base, gaining a proton to become H_2CO_3 .

23. **D**

You want to change the hydrogen ion concentration from 0.1 M (pH of 1) to 0.01 M (pH of 2).

The HCl is completely dissociated, so the number of moles of H^+ will remain constant as we dilute the solution.

Moles = (molarity)(volume) = Constant

$$(M_1)(V_1) = (M_2)(V_2)$$

$$(0.1\text{ M})(1\text{ L}) = (0.01\text{ M})(V_2)$$

So, $V_2 = 10\text{ L}$, which means that 9 L must be added.

24. **A**

A pH of 5 means that $[\text{H}^+] = 1 \times 10^{-5}$.

$$K_a = \frac{[\text{H}^+][\text{A}^-]}{[\text{HA}]}$$

For every HA that dissociates, you get one H^+ and one A^- , so $[\text{H}^+] = [\text{A}^-] = 1 \times 10^{-5}$.

The acid is weak, so you can assume that very little HA dissociates and that the concentration of HA remains 1-molar.

$$K_a = \frac{[\text{H}^+][\text{A}^-]}{[\text{HA}]} = \frac{(1 \times 10^{-5})(1 \times 10^{-5})}{(1)} = 1 \times 10^{-10}.$$

25. **A**

For conjugates, $(K_a)(K_b) = K_w = 1 \times 10^{-14}$

$$K_b = \frac{K_w}{K_a} = \frac{(1 \times 10^{-14})}{(1 \times 10^{-2})} = 1 \times 10^{-12}$$

26. **B**

The reaction here is $\text{NH}_3 + \text{H}^+ \rightleftharpoons \text{NH}_4^+$. At equilibrium, the moles of NH_3 and H^+ would be equal, leaving behind NH_4^+ ions, which will then donate ions to water, creating an acidic medium.

27. **D**

At the equivalence point, the moles of acid are equal to the moles of base.

Moles base = $(1.0 \text{ M})(0.030 \text{ L}) = 0.030 \text{ mol base} = 0.030 \text{ moles acid}$

It requires 15.0 mL of acid to reach equivalence, so:

$$\frac{0.030 \text{ mol}}{0.015 \text{ L}} = 2.0 \text{ M}$$

28. **A**

The reaction occurring is $\text{NH}_3 + \text{H}^+ \rightleftharpoons \text{NH}_4^+$. During the first buffer region, all added hydrogen ions immediately react with NH_3 to create NH_4^+ . NH_3 remains in excess until equilibrium is achieved.

29. **A**

There are no sulfate ions in solution, which means that whatever the sulfate was attached to was the limiting reactant. Thus, silver must have been attached to the excess reactant, which is AgNO_3 . (Remember, Ag_2SO_4 was the product, not a reactant.)

30. **D**

Leaving the container uncovered will cause some of the water molecules to evaporate. This allows some Ag^+ and SO_4^{2-} ions to “fall” out of solution and combine to create more Ag_2SO_4 .

31. **B**

For a precipitate to form, $Q > K_{sp}$. In this case, $Q = [\text{Ag}^+]^2[\text{SO}_4^{2-}]$. The concentrations in (B) lead to a Q of 1.0×10^{-4} , which is greater than the given K_{sp} of 1.0×10^{-5} . The other options lead to a Q value that is equal to or less than K_{sp} .

32. **A**

The reaction here is $\text{Pb (s)} + \text{Cu}^{2+} \rightleftharpoons \text{Cu (s)} + \text{Pb}^{2+}$. Increasing the concentration of Pb^{2+} causes Q to increase, and as all voltaic cells have $K > 1$, that brings Q closer to K . For any voltaic cell, the voltage is zero at equilibrium, so bringing the cell closer to equilibrium brings the voltage closer to zero.

33. **B**

Percent dissociation is inversely proportional to concentration. Adding some water to the solution will

cause the concentration of the HF to decrease, leading to a greater percent dissociation.

34. **D**

$\text{H}_2\text{O} \rightleftharpoons \text{H}^+ + \text{OH}^-$ is an endothermic process, requiring energy in order for the bonds within the water molecules to break. Thus, heat is a reactant, and as temperature increases the reaction will shift right. Because this is a temperature shift, this causes a permanent increase in the value of K_w . This, in turn, causes the concentrations of H^+ and OH^- to increase, and the $-\log$ of an increased concentration leads to a decreased value for pH.

35. **D**

Via $K_{eq} = e^{-\frac{\Delta G}{RT}}$, if ΔG is negative the value for K will be greater than one. Via $\Delta G = \Delta H - T\Delta S$, ΔG is always negative when ΔH is negative and ΔS is positive.

36. **D**

The weaker the O–H bond is in an oxoacid, the stronger the acid will be, because the H^+ ions are more likely to dissociate. The O–F bond in HOF is stronger than the O–Cl bond in HOCl because fluorine is smaller (and thus more electronegative) than chlorine. If the O–F bond is stronger, the O–H bond is correspondingly weaker, making HOF the stronger acid.

37. **A**

A buffer is made up of either a weak acid and its salt or a weak base and its salt. Choice (B) has a strong acid and

strong base, (C) has a strong acid and its salt, and (D) has a weak acid and a weak base.

38. **B**

Increasing the pressure in an equilibrium reaction with any gas molecules causes a shift to the side with fewer gas molecules—in this case, the product.

39. **C**

The strength of an acid is dependent on the amount it dissociates in solution. A low dissociation is signified by a low presence of hydrogen ions. The weakest acid is 3, (C).

40. **C**

Use $M_1V_1 = M_2V_2$ to determine the necessary volume of stock solution.

$$(18.0\text{ M})V_1 = (3.0\text{ M})(1.0\text{ L})$$

$$V_1 = 0.167\text{ L} = 167\text{ mL}$$

When creating solutions with acid, you always add some water first, as the process is extremely exothermic and the water will absorb the generated heat.

41. **D**

Adding (or removing) any species in an equilibrium reaction does not change the equilibrium constant and also does not change the magnitude of the Gibbs free energy at standard conditions.

Free-Response

1. (a) The solubility product is the same as the equilibrium expression, but because the reactant is a solid, there is no denominator.

$$K_{sp} = [\text{Ca}^{2+}][\text{OH}^-]^2$$

- (b) Use the solubility product.

$$K_{sp} = [\text{Ca}^{2+}][\text{OH}^-]^2$$

$$5.5 \times 10^{-6} = (x)(2x)^2 = 4x^3$$

$$x = 0.01 \text{ M for } \text{Ca}^{2+}$$

One mole of calcium hydroxide produces 1 mole of Ca^{2+} , so the concentration of $\text{Ca}(\text{OH})_2$ must be 0.01 M.

$$\text{Moles} = (\text{molarity})(\text{volume})$$

$$\text{Moles of } \text{Ca}(\text{OH})_2 = (0.01 \text{ M})(0.500 \text{ L}) = 0.005 \text{ moles}$$

$$\text{Grams} = (\text{moles})(\text{MW})$$

$$\text{Grams of } \text{Ca}(\text{OH})_2 = (0.005 \text{ mol})(74 \text{ g/mol}) = 0.37 \text{ g}$$

- (c) You can find $[\text{OH}^-]$ from (b).

If $[\text{Ca}^{2+}] = 0.01 \text{ M}$, then $[\text{OH}^-]$ must be twice that, so $[\text{OH}^-] = 0.02 \text{ M}$.

$$\text{pOH} = -\log[\text{OH}^-] = 1.7$$

$$\text{pH} = 14 - \text{pOH} = 14 - 1.7 = 12.3$$

- (d) Find the new $[\text{OH}^-]$. The hydroxide already present is small enough to ignore, so use only the hydroxide just

added.

$$\text{Molarity} = \frac{\text{moles}}{\text{liters}}$$

$$[\text{OH}^-] = \frac{(1.0 \text{ mol})}{(0.500 \text{ L})} = 2.0 \text{ M}$$

$$[\text{OH}^-] = 2.0 \text{ M}$$

Now use the K_{sp} expression.

$$K_{sp} = [\text{Ca}^{2+}][\text{OH}^-]^2$$

$$5.5 \times 10^{-6} = [\text{Ca}^{2+}](2.0 \text{ M})^2$$

$$[\text{Ca}^{2+}] = 1.4 \times 10^{-6} \text{ M}$$

2. (a) Entropy increases when a salt dissociates because aqueous particles have more randomness than a solid.
- (b) Most salt solution processes are endothermic, and endothermic processes are favored by an increase in temperature. Therefore, increasing temperature will increase the solubility of most salts.
- (c) In an endothermic reaction, the energy is part of the reactants. Increasing the temperature would thus shift the reaction to the right.
- (d) The solution was already saturated, which means $[\text{Na}^+]$ and $[\text{Cl}^-]$ were already at their maximum possible values. If the volume of water decreases, the number of moles of Na^+ and Cl^- present in solution must also decrease in order for their concentrations to remain constant. In order for that to happen, some of the sodium and chloride ions present

will form a precipitate, increasing the mass of NaCl(s) present.

3. A student tests the conductivity of three different acid samples, each with a concentration of 0.10 M and a volume of 20.0 mL . The conductivity was recorded in microsiemens per centimeter in the table below:

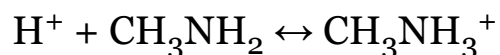
Sample	Conductivity ($\mu\text{S/cm}$)
1	26,820
2	8655
3	35,120

- (a) The three acids are known to be HCl , H_2SO_4 , and H_3PO_4 . Identify which sample is which acid. Justify your answer.

H_3PO_4 is the only weak acid, meaning it will not dissociate completely in solution. Therefore, it is acid 2. H_2SO_4 and HCl are both strong acids that will dissociate completely, but H_2SO_4 is diprotic, meaning there will be more H^+ ions in solution and thus a higher conductivity. Therefore, H_2SO_4 is acid 3, and HCl is acid 1.

- (b) The HCl solution is then titrated with a 0.150 M solution of the weak base methylamine, CH_3NH_2 . ($K_b = 4.38 \times 10^{-4}$).

- (i) Write out the net ionic equation for this reaction.



- (ii) Determine the pH of the solution after 20.0 mL of methylamine has been added.

First, the number of moles of hydrogen ions and methylamine need to be determined.

$$\text{H}^+: 0.10 \text{ M} = \frac{n}{0.020 \text{ L}} \quad n = 0.0020 \text{ mol}$$

$$\text{CH}_3\text{NH}_2: 0.150 \text{ M} = \frac{n}{0.020 \text{ L}} \quad n = 0.0030 \text{ mol}$$



I	0.0020	0.0030	0
C	-0.0020	-0.0020	+0.0020
E	0	0.0010	0.0020

The new volume of the solution is 40.0 mL, which was used to calculate the concentrations of the ions in solution at equilibrium:

$$[\text{CH}_3\text{NH}_2] = \frac{0.0010 \text{ mol}}{0.040 \text{ L}} = 0.025 \text{ M}$$

$$[\text{CH}_3\text{NH}_3^+] = \frac{0.0020 \text{ mol}}{0.040 \text{ L}} = 0.050 \text{ M}$$

Finally, use the Henderson-Hasselbalch equation.

$$\text{pOH} = \text{p}K_{\text{b}} + \log \frac{[\text{CH}_3\text{NH}_3^+]}{[\text{CH}_3\text{NH}_2]}$$

$$\text{pOH} = -\log(4.38 \times 10^{-4}) + \log \frac{0.050}{0.025}$$

$$\text{pOH} = 3.36 + 0.30 = 3.66$$

$$\text{pH} + \text{pOH} = 14$$

$$\text{pH} + 3.66 = 14$$

$$\text{pH} = 10.34$$

$$4. \quad (a) \quad K_c = \frac{[\text{NH}_3]^2}{[\text{N}_2][\text{H}_2]^3}$$

$$K_c = \frac{(0.20)^2}{(0.50)(0.20)^3} = 10$$

$$(b) \quad \Delta G = -RT \ln K$$

$$\Delta G = -(8.31 \text{ J/mol} \times \text{K})(570 \text{ K}) \ln (10)$$

$$\Delta G = (-4740)(2.3)$$

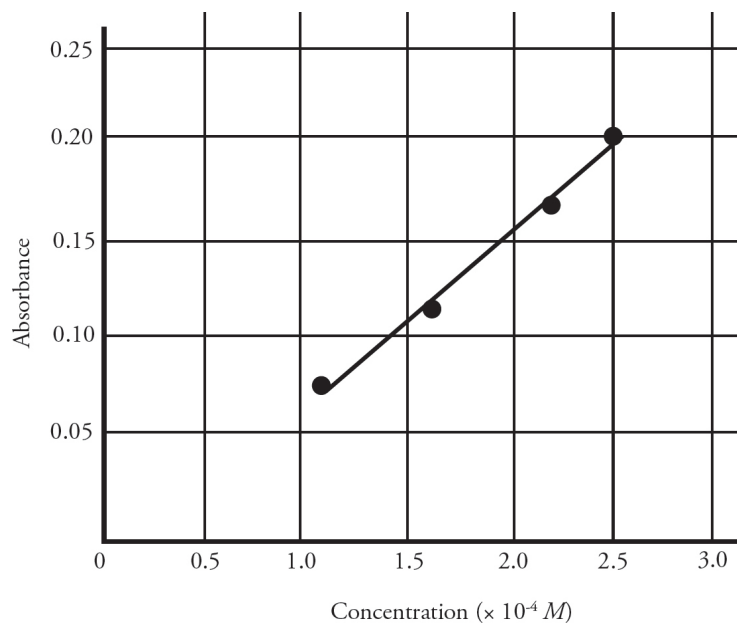
$$\Delta G = -11,000 \text{ J/mol}$$

(c) (i) An increase in temperature will shift the reaction to the left, causing an increase in the H_2 concentration in the short term. The temperature increase also causes a decrease in the value for the equilibrium constant. A smaller equilibrium constant means the species in the denominator of the mass action expression will be present in increased concentrations at equilibrium, so $[\text{H}_2]$ will remain increased.

(ii) An increase in volume favors the direction that produces more moles of gas. In this case, that's the reverse direction, so the concentration of H_2 will increase in the short term. However, an increased chamber volume will eventually reduce the concentrations of all species at equilibrium, as the number of moles remains constant and concentration is calculated using moles/volume. Thus, $[\text{H}_2]$ will decrease by the time equilibrium re-establishes.

- (iii) Adding N_2 will initially shift the reaction right, and the concentration of H_2 will decrease. However, by the time equilibrium is re-established, there will be more moles of all three substances in the container, so ultimately $[\text{H}_2]$ will be increased.
- (iv) The addition of He gas at a constant volume will increase the total pressure in the container, but the partial pressure of each species will remain unchanged. Thus, $[\text{H}_2]$ will be the same. Note that if the He were added at constant pressure, that would lead to a different result, as described on [this page](#).

5. (a) (i)



(ii) Slope-intercept form is $y = mx + b$. So:

$$y = 879x - 0.0240$$

It is worth noting that in a perfect world, the y -intercept for this graph would be zero; that is, when there is no FeSCN^{2+} present the solution will not absorb light. Due

to the nature of experimental data in the real world being not perfect, though, it is best in this case to include the approximate y -intercept in your best-fit line, as it may account for variations in your equipment.

(b) (i) Using the equation from part (a):

$$y = 879x - 0.0240$$

$$0.134 = 879x - 0.0240$$

$$x = 1.80 \times 10^{-4} M$$

(ii) An ICE chart will be very helpful here. To do that, you first need to calculate the initial concentration of the Fe^{3+} and the SCN^- . They both start with the same volumes and concentrations, so one calculation will suffice. Start with calculating the moles present:

$$0.0025 M = \frac{n}{0.0050 \text{ L}} \quad n = 1.3 \times 10^{-5} \text{ mol}$$

Then, you have to divide by the total volume of both solutions combined (10.0 mL) in order to find out the concentration of each ion.

$$\frac{1.3 \times 10^{-5} \text{ mol}}{0.010 \text{ L}} = 1.3 \times 10^{-3} M$$

The ICE chart thus looks like this:



I	1.3×10^{-3}	1.3×10^{-3}	0
C	-1.80×10^{-4}	-1.80×10^{-4}	$+1.80 \times 10^{-4}$
E	1.1×10^{-3}	1.1×10^{-3}	1.80×10^{-4}

So at equilibrium $[\text{Fe}^{3+}] = [\text{SCN}^-] = 1.1 \times 10^{-3} M$.

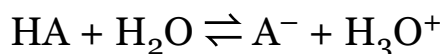
$$(iii) K_{eq} = \frac{[FeSCN^{2+}]}{[Fe^{3+}][SCN^{-}]} = \frac{1.80 \times 10^{-4}}{(1.1 \times 10^{-3})(1.1 \times 10^{-3})} = 150$$

- (c) (i) Changing the temperature of a reaction is the only way to change the equilibrium constant, which is what is happening here. As the solution becomes lighter, the amount of $FeSCN^{2+}$ decreases, which indicates a shift to the left. Doing so causes there to be more products and fewer reactants at equilibrium, which lowers the value of the equilibrium constant.
- (ii) Using the trick for temperature-caused shifts, you can see that a shift to the left means that there must have been heat on the products side of the equilibrium. This means the reaction is exothermic.
6. (a) (i) Between these volumes lies the buffer region of the titration, in which both $[HA]$ and $[A^{-}]$ contribute significantly to the pH of the solution.
- (ii) At equivalence, the only species present which affects the pH of the solution is the conjugate base, A^{-} .
- (iii) In this region, the strong base is in excess, and $[OH^{-}]$ determines the solution.
- (b) (i) As the reaction progress, OH^{-} will take protons from HA to create A^{-} . There will be more HA in solution until half of it has been deprotonated. The point at which that occurs is the half-equivalence point at 7.50 mL. So, $[HA] > [A^{-}]$ from 0 to 7.5 mL.
- (ii) The only time these values are equal is when exactly half of the HA has been converted to A^{-} , which occurs at 7.50 mL.

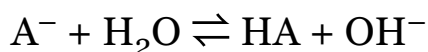
(iii) For the remainder of the titration (7.50 mL+), there will be more conjugate base present than there will be of original weak acid.

(c) (i) and (ii)

Neither of these values will ever be equal to zero. Prior to the titration beginning, there is some A^- in solution via the reaction of HA with water, as follows:



Adding OH^- through titrating the strong base in will cause the HA molecules present to deprotonate fully, but the conjugate base that is created will, in turn, also react with water to create additional HA molecules:



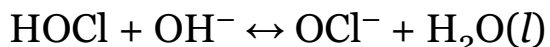
As both the reaction of the weak acid with water and that of its conjugate base with water will always be occurring, there will always be some of each species in solution.

(d)

- Twice as much OH^- is being added per drop, so the pH changes charted will occur over only 15.0 mL instead of 30.0 mL (for instance, equivalence will occur at 7.50 mL instead of 15.0 mL).
- The final pH of the solution will be slightly higher, as a higher concentration of NaOH will lead to a higher pH in the region where $[OH^-]$ dictates the pH of the solution.

7. (a) The hypochlorous acid will donate its proton to the hydroxide. The sodium ions are spectators and would not

appear in the net ionic equation.



- (b) At the equivalence point, the moles of acid are equal to the moles of base.

$$\text{Moles of base} = (1.47 \text{ M})(0.03423 \text{ L}) = 0.0503 \text{ mol}$$

$$\text{Concentration acid} = (0.0503 \text{ mol}/0.02500 \text{ L}) = 2.01 \text{ M}$$

- (c) This calls for an ICE chart. You will first need the moles of both the acid and base.

$$\text{Moles HOCl} = (2.01 \text{ M})(0.02500 \text{ L}) = 0.0503 \text{ mol HOCl}$$

$$\text{Moles OH}^- = (1.47 \text{ M})(0.02855 \text{ L}) = 0.0420 \text{ mol OH}^-$$

Putting those numbers into the ICE chart:

	HOCl	OH ⁻	OCl ⁻	H ₂ O
Initial	0.0503	0.0420	0	X
Change	-0.0420	-0.0420	0.0420	X
Equilibrium	0.0083	0	0.0420	X

Now that you know the number of moles at equilibrium of each species, you need to determine their new concentrations. The total volume of the solution at equilibrium is 25.00 mL + 28.55 mL = 53.55 mL. So the concentrations at equilibrium will be:

$$[\text{HOCl}] = 0.0083 \text{ mol}/0.05355 \text{ L} = 0.15 \text{ M}$$

$$[\text{OCl}^-] = 0.0420 \text{ mol}/0.05355 \text{ L} = 0.784 \text{ M}$$

To finish off, use Henderson-Hasselbalch:

$$\text{pH} = \text{p}K_a + \log \frac{[\text{OCl}^-]}{[\text{HOCl}]}$$
$$\text{pH} = -\log (3.5 \times 10^{-8}) + \log \frac{(0.784)}{(0.15)}$$

$$\text{pH} = 7.5 + \log (5.2)$$

$$\text{pH} = 7.5 + 0.72$$

$$\text{pH} = 8.2$$

- (d) (i) If additional NaOH is added, that would mean a larger number of moles of NaOH added to the flask, and thus more apparent moles of HOCl would be present at the equivalence point. More apparent moles of HOCl would lead to a larger numerator in the acid molarity calculation and thus a larger apparent molarity. As the student's calculated molarity was too low, this could not have caused the error.
- (ii) If the buret was not rinsed with the NaOH prior to filling it, the concentration of the NaOH would be diluted by the water inside the buret. The student would then have to add a greater volume of NaOH to reach equivalence. However, in the calculations, using the original concentration of NaOH multiplied by the higher volume would lead to an artificially high number of moles of NaOH and thus, more apparent moles HOCl at equivalence. More apparent moles of HOCl would lead to a larger numerator in the acid molarity calculation and thus a larger apparent molarity. As the student's calculated molarity was too low, this could not have caused the error.
- (iii) If the student did not add enough acid to the flask, that would cause the denominator in the molarity of the acid

calculation to be artificially high. This, in turn, would make the calculated acid molarity be artificially low. This matches up with the student's results and could be an acceptable source of error.

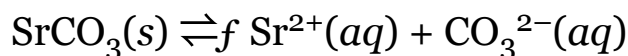
8. A student is tasked with determining the identity of an unknown carbonate compound with a mass of 1.89 g. The compound is first placed in water, where it dissolves completely. The K_{sp} values for several carbonate-containing compounds are given below.

Compound	K_{sp}
Lithium carbonate	8.15×10^{-4}
Nickel (II) carbonate	1.42×10^{-7}
Strontium carbonate	5.60×10^{-10}

- (a) In order to precipitate the maximum amount of the carbonate ions from solution, which of the following should be added to the carbonate solution: LiNO_3 , $\text{Ni}(\text{NO}_3)_2$, or $\text{Sr}(\text{NO}_3)_2$? Justify your answer.

The student should use the strontium nitrate. Using it would create strontium carbonate, which has the lowest K_{sp} value. That means it is the least soluble carbonate compound of the three and will precipitate the most possible carbonate ions out of solution.

- (b) For the carbonate compound that contains the cation chosen in part (a), determine the concentration of each ion of that compound in solution at equilibrium.



$$K_{\text{sp}} = [\text{Sr}^{2+}][\text{CO}_3^{2-}]$$

$$5.60 \times 10^{-10} = (x)(x)$$

$$x = 2.37 \times 10^{-5} \text{ M} = [\text{Sr}^{2+}] = [\text{CO}_3^{2-}]$$

- (c) When mixing the solution, should the student ensure the carbonate solution or the nitrate solution is in excess? Justify your answer.

The nitrate solution should be in excess. In order to create the maximum amount of precipitate, enough strontium ions need to be added to react with all of the carbonate ions originally in solution. Having excess strontium ions in solution after the precipitate forms will not affect the calculated mass of the carbonate in the original sample.

- (d) After titrating sufficient solution to precipitate out all of the carbonate ions, the student filters the solution before placing it in a crucible and heating it to drive off the water. After several heatings, the final mass of the precipitate remains constant and is determined to be 2.02 g.

- (i) Determine the number of moles of precipitate.

$$2.02 \text{ g SrCO}_3 \times \frac{1 \text{ mol SrCO}_3}{147.63 \text{ g SrCO}_3} = 1.37 \times 10^{-2} \text{ mol SrCO}_3$$

- (ii) Determine the mass of carbonate present in the precipitate.

$$1.37 \times 10^{-2} \text{ mol SrCO}_3 \times \frac{1 \text{ mol CO}_3^{2-}}{1 \text{ mol SrCO}_3} \times \frac{60.01 \text{ g CO}_3^{2-}}{1 \text{ mol CO}_3^{2-}} = 0.822 \text{ g CO}_3^{2-}$$

- (e) Determine the percent, by mass, of carbonate in the original sample.

$$\frac{0.82 \text{ g}}{1.89 \text{ g}} \times 100 = 43.4\% \text{ CO}_3^{2-}$$

- (f) Is the original compound most likely lithium carbonate, sodium carbonate, or potassium carbonate? Justify your answer.

The mass percent of carbonate in each compound must be compared to the experimentally determined mass percent of carbonate in the sample.

$$\text{Li}_2\text{CO}_3: \frac{60.01}{73.89} \times 100 = 81.2\%$$

$$\text{Na}_2\text{CO}_3: \frac{60.01}{105.99} \times 100 = 56.6\%$$

$$\text{K}_2\text{CO}_3: \frac{60.01}{138.21} \times 100 = 43.4\%$$

The compound is most likely potassium carbonate.

Chapter 9

Laboratory Overview

INTRODUCTION

The AP Chemistry Exam will test your knowledge of basic lab techniques, as well as your understanding of accuracy and precision and your ability to analyze potential sources of error in a lab. In this section, we will discuss safety and accuracy precautions, laboratory equipment, and laboratory procedures.

SAFETY



Here are some basic safety rules that might turn up in test questions.

- Don't put chemicals in your mouth. You were told this when you were four years old, and it still holds true for the AP Chemistry Exam.
- When diluting an acid, always add the acid to the water. This is to avoid the spattering of hot solution.
- Always work with good ventilation; many common chemicals are toxic.
- When heating substances, do it slowly. When you heat things too quickly, they can spatter, burn, or explode.

ACCURACY

Here are some rules for ensuring the accuracy of experimental results.

- When titrating, rinse the buret with the solution to be used in the titration instead of with water. If you rinse the buret with water, you might dilute the solution, which will cause the volume added from the buret to be too large.

- Allow hot objects to return to room temperature before weighing. Hot objects on a scale create convection currents that may make the object seem lighter than it is.
- Don't weigh reagents directly on a scale. Use a glass or porcelain container to prevent corrosion of the balance pan.
- When collecting a gas over water, remember to take into account the pressure and volume of the water vapor.
- Don't contaminate your chemicals. Never insert another piece of equipment into a bottle containing a chemical. Instead you should always pour the chemical into another clean container. Also, don't let the inside of the stopper for a bottle containing a chemical touch another surface.
- When mixing chemicals, stir slowly to ensure even distribution.
- Be conscious of significant figures when you record your results. The number of significant figures you use should indicate the accuracy of your results.
- Be aware of the difference between accuracy and precision. A measurement is accurate if it is close to the accepted value. A series of measurements is precise if the values of all of the measurements are close together.

SIGNIFICANT FIGURES

When taking measurements in lab, your measurement will always have a certain number of significant figures. For instance, if you are using a balance to measure the weight of an object to the hundredths place, you might get 23.15 g. That number has four significant figures. The balance is no more accurate than that; you can not say the mass of the object is 23.15224 g.

It's important to be able to identify the number of significant figures in any number given to you on the AP Exam. There's a plethora of information on ways to count significant figures out there, but we've reduced it to two simple rules.

1. For numbers without a decimal place, you count every number except for trailing zeroes (those which appear after all non-zeroes). So, 100 mL only has one significant figure, and 250 mL has two. On the other hand, 105 g has three significant figures—the zero in that measurement does not trail all other numbers.
2. For numbers with a decimal place, you count every number except for leading zeroes (those which appear before all non-zeroes). The number 0.052 has 2 significant figures, but 0.0520 g would have three (trailing zeroes DO count in numbers with decimals). Most (but not all) values you get on the AP Exam will have decimal places.

This leads to numbers that otherwise might look very strange. For instance, it is not unusual to see numbers with a decimal at the end but no numbers past it. That's because a number like 100. g has three significant figures, but 100 g only has one. In science, 100. g is not the same as 100 g, which is also not the same as 100.0 g. All of those values have different numbers of significant figures, and that implies different levels of accuracy. Significant figures only apply to measurements with units; pure unitless numbers (such as those you find in math) do not follow these rules. Nonetheless, many math books do have units on practice problems and then ignore significant figures. Yes, it's wrong to do that, but don't tell your math teacher.



More Great Books

If all this math talk is worrying you, check out these other books from The Princeton Review to help you in your math review:

High School Geometry Unlocked

High School Algebra I Unlocked

High School Algebra II Unlocked

When doing calculations, it is important that any calculated value cannot be more accurate than the measurements used in the calculation. Essentially, significant figures can tell you how to round your answers correctly. Again, this can be divided into two categories:

1. When multiplying and dividing, your answer cannot have more significant figures than your least accurate measurement. For instance:

$$2.50 \text{ g} / 12 \text{ cm}^3 = 0.20833 \text{ g/cm}^3$$

is wrong. Your two measurements have three significant figures and two significant figures, respectively. Your answer cannot have five. It has to have only two because that's how many figures your least accurate measurement had. The correct answer is 0.21 g/cm^3 .

2. When adding and subtracting, your answer cannot have more figures after the decimal place than your value with the least number of figures after its decimal place. For instance:

$$1.435 \text{ cm} + 12.1 \text{ cm} = 13.535 \text{ cm}$$

is wrong. Your values have three figures after the decimal and one figure after the decimal, respectively. Your answer cannot have more than one figure past the decimal, because that's how many figures your least accurate measurement had. The correct answer is 13.5 cm.

On a side note, counting numbers are considered to have an infinite number of significant figures. If you are doing a molar mass conversion and you identify the conversion as 22.99 g of sodium = 1 mol of sodium, your eventual answer will have four significant figures. That 1 mol represents *exactly* 1 mol of sodium, which is essentially a 1 with an infinite number of zeroes past the decimal place. This will usually come up when dealing with stoichiometry—the coefficients in a balanced equation are counting numbers of moles and don't count when considering the number of significant figures your answer should have.

You should get into the habit of making sure your calculations always have the correct number of significant figures as you do them, both in labs and on any practice problems you do in class. The AP Exam may have questions on significant figures in a lab, and they are typically integrated into the free-response section. Additionally, you may lose points for the incorrect number of significant figures when you do calculations. Along with making sure you have the correct units on your number, making sure you do your calculations to the correct number of significant figures is a good laboratory practice.

EXPERIMENTAL DESIGN

The AP Chemistry Exam is heavily focused on students being able to understand the various types of experiments that can be undertaken in order to reinforce the concepts that are taught in the classroom. Some, if not most, of the free-response questions will be based on laboratory investigations. Here are the seven types that, based on the current framework for the exam, you should be familiar with. We've listed the test-maker's expectations (in their own words), a brief

description of the experiment, and where in this book you can find additional explanations and practice problems.

Spectrophotometry

LO 1.16: The student can design and/or interpret the results of an experiment regarding the absorption of light to determine the concentration of an absorbing species in a solution.

Spectrophotometry (aka colorimetry) is often used to measure the concentration of colored solutions. This type of experiment is often done in relation with reaction rates and/or equilibrium calculations.

Conceptual Explanation: [this page](#)

Free-Response Examples: Chapter 6 FRQ #8, Chapter 8 FRQ #5

Gravimetric Analysis

LO 1.19: The student can design, and/or interpret data from, an experiment that uses gravimetric analysis to determine the concentration of an analyte in a solution.

Gravimetric analysis is intrinsically tied in with precipitation reactions and mass percent calculations.

Conceptual Explanation: [this page](#)

Free-Response Examples: Chapter 5 FRQ #1, Chapter 8 FRQ #8

Acid-Base Titrations

LO 1.20: The student can design, and/or interpret data from, an experiment that uses titration to determine the concentration of an analyte in a solution.

Acid-base titrations are by far the most common form of this learning objective that you will see on the exam. Acid-base titrations

typically use indicators to cause the color to change at or near the equivalence point.

Conceptual Explanation: Pages 286–287

Free-Response Examples: Chapter 8 FRQs #6 and #7

Component Separation

LO 2.10: The student can design and/or interpret the results of a separation experiment (filtration, paper chromatography, column chromatography, or distillation) in terms of the relative strength of interactions among and between the components.

Understanding the best way to separate unique substances out of a mixture ties directly in with concepts of intermolecular forces and bonding. Out of the seven types of experiments, this one is generally the least mathematical and the most conceptual.

Conceptual Explanation: [this page](#)

Free-Response Examples: Chapter 4 FRQs #1 and #4

Redox Titrations

LO 3.9: The student is able to design and/or interpret the results of an experiment involving a redox titration.

As the name indicates, these are titrations that are focused on oxidation-reduction reactions. These do not always use indicators (although they can), but instead some of the solutions involved are colored themselves.

Conceptual Explanation: [this page](#)

Free-Response Examples: Chapter 5 FRQs #2 and #7

Kinetics

LO 4.1: The student is able to design and/or interpret the results of an experiment regarding the factors (i.e., temperature, concentration, surface area) that may influence the rate of a reaction.

Determining the rate of reaction can usually only be done experimentally. As such, that makes kinetics an easy topic to study via experimental design. Graphical analysis is particularly common in these experiments.

Conceptual Explanation: [this page](#)

Free-Response Examples: Chapter 6 FRQs #4 and #8

Calorimetry

LO 5.7 The student is able to design and/or interpret the results of an experiment in which calorimetry is used to determine the change in enthalpy of a chemical process (heating/cooling, phase transition, or chemical reaction) at constant pressure.

The most common experimental data you will deal with involving any kind of thermodynamics calculations begins with calorimetry. Most thermodynamic data is gathered from existing data tables/research, but calorimetry is something that any high school student can experience during lab.

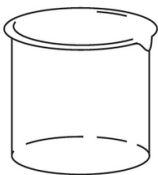
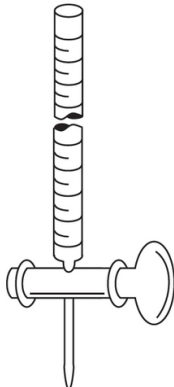
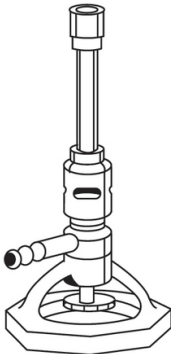
Conceptual Explanation: [this page](#)

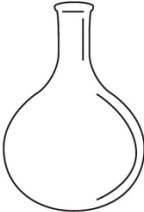
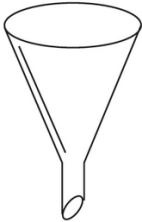
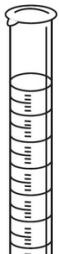
Free-Response Example: Chapter 7 FRQ #6

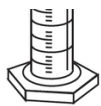
There will certainly be other types of experiments tested on the AP Exam—there are a countless number of ways to set up and design experiments to test various chemistry concepts. However, focusing on these types is an excellent place to start.

LABORATORY EQUIPMENT

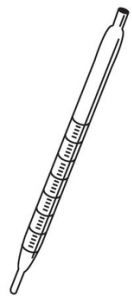
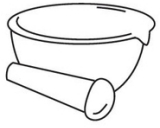
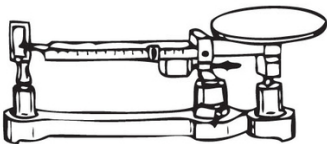
The following charts show some standard chemistry lab equipment.

 Beaker	Used to hold and pour liquids—also your favorite muppet.
 Buret	Used to add small but precisely measured volumes of liquid to a solution. Burets are used frequently in titration experiments.
 Burner	Used to apply heat and to wake up sleeping AP Chemistry students.
 Crucible tongs	Used to handle objects that are too hot to touch (careful though, they can break test tubes!).
 Dropper pipette	Used to add small amounts of liquid to a solution, but only when a precise volume is not needed, because the drops themselves should be consistent only for one particular dropper (for example, adding an indicator, comparing the amount of drops of strong acid or base needed for a pH change in different solutions).

 Erlenmeyer flask	A flask used for heating liquids. The conic shape allows stirring.
 Evaporating dish	Used to hold liquids for evaporation. The wide mouth allows vapor to escape.
 Florence flask	Used for boiling of liquids. The small neck prevents excessive evaporation and splashing.
 Forceps	A fancy name for tweezers.
 Funnel	Used to get liquids into a smaller container. Doubles as a cute hat when inverted.
	Used for measuring precisely a volume of liquid to be poured all at once (rather than dripped from a buret).



Graduated
cylinder

 Graduated pipette	Used to transfer small and precise volumes of liquid from one container to another (the gradations indicate the volume).
 Metal spatula	Used to scoop and transport powders.
 Mortar and pestle	Used to grind solids into powders suitable for dissolving or mixing.
 Pipette bulb	Rubber bulb used to draw liquid into pipette. In a pinch, could be used as a miniature clown nose.
 Platform balance (triple beam)	A very precise scale operated by moving a set of three weights (typically corresponding to 100, 10, and 1 gram increments). A measurement will proceed like this: Rear weight is in the notch reading 30 g. Middle weight is in the notch reading 200 g. Front beam weight reads 3.86 g. The sample weighs $200 + 30 + 3.86 = 233.86$ g.
 Ring clamp	Used to hold funnels or other vessels in conjunction with a stand.
	A hard tipped rubber scraper used to transfer precipitate.



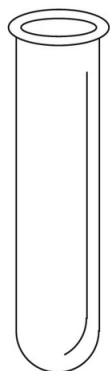
Rubber
policeman



Safety goggles

Used by chemists to protect their eyes during all laboratory experiments.

Many students like to wear these on their foreheads or as a headband, but they really must be worn over your eyes to protect you.



Test tube

Used to contain samples, especially when heating. Sometimes babies come out of them.



Thermometer

Measures temperature of a solution (don't use these to measure your body temperature, you don't know where they've been).



Volumetric
pipette

Used to transfer small amounts of liquid. The big difference between a graduated pipette and a volumetric pipette is that the volumetric type is suited for only one particular volume. Because of this, they are extremely accurate.

Part VI

Practice Test 2

[Click here](#) to download and print the PDF version of this exercise.

AP[®] Chemistry Exam

SECTION I: Multiple-Choice Questions

DO NOT OPEN THIS BOOKLET UNTIL YOU ARE TOLD TO DO SO.

At a Glance

Total Time

1 hour and 30 minutes

Number of Questions

60

Percent of Total Grade

50%

Writing Instrument

Pencil required

Instructions

Section I of this examination contains 60 multiple-choice questions. Fill in only the ovals for numbers 1 through 60 on your answer sheet.

CALCULATORS MAY NOT BE USED IN THIS PART OF THE EXAMINATION.

Indicate all of your answers to the multiple-choice questions on the answer sheet. No credit will be given for anything written in this exam booklet, but you may use the booklet for notes or scratch work. After you have decided which of the suggested answers is best, completely fill in the corresponding oval on the answer sheet. Give only one answer to each question. If you change an answer, be sure that the previous mark is erased completely. Here is a sample question and answer.

Sample Question

Chicago is a

- (A) state
- (B) city
- (C) country
- (D) continent

Sample Answer

☒ (A) ☐ (B) ☐ (C) ☐ (D)

Use your time effectively, working as quickly as you can without losing accuracy. Do not spend too much time on any one question. Go on to other questions and come back to the ones you have not answered if you have time. It is not expected that everyone will know the answers to all the multiple-choice questions.

About Guessing

Many candidates wonder whether or not to guess the answers to questions about which they are not certain. Multiple-choice scores are based on the number of questions answered correctly. Points are not deducted for incorrect answers, and no points are awarded for unanswered questions. Because points are not deducted for incorrect answers, you are encouraged to answer all multiple-choice questions.

On any questions you do not know the answer to, you should eliminate as many choices as you can, and then select the best answer among the remaining choices.

CHEMISTRY

SECTION I

Time—1 hour and 30 minutes

INFORMATION IN THE TABLE BELOW AND ON THE FOLLOWING PAGES MAY BE USEFUL IN ANSWERING THE QUESTIONS IN THIS SECTION OF THE EXAMINATION.

DO NOT DETACH FROM BOOK.

PERIODIC TABLE OF THE ELEMENTS

1 H 1.008																	2 He 4.00
3 Li 6.94	4 Be 9.01											5 B 10.81	6 C 12.01	7 N 14.01	8 O 16.00	9 F 19.00	10 Ne 20.18
11 Na 22.99	12 Mg 24.30											13 Al 26.98	14 Si 28.09	15 P 30.97	16 S 32.06	17 Cl 35.45	18 Ar 39.95
19 K 39.10	20 Ca 40.08	21 Sc 44.96	22 Ti 47.90	23 V 50.94	24 Cr 52.00	25 Mn 54.94	26 Fe 55.85	27 Co 58.93	28 Ni 58.69	29 Cu 63.55	30 Zn 65.39	31 Ga 69.72	32 Ge 72.59	33 As 74.92	34 Se 78.96	35 Br 79.90	36 Kr 83.80
37 Rb 85.47	38 Sr 87.62	39 Y 88.91	40 Zr 91.22	41 Nb 92.91	42 Mo 95.94	43 Tc (98)	44 Ru 101.1	45 Rh 102.91	46 Pd 106.42	47 Ag 107.87	48 Cd 112.41	49 In 114.82	50 Sn 118.71	51 Sb 121.75	52 Te 127.60	53 I 126.91	54 Xe 131.29
55 Cs 132.91	56 Ba 137.33	57 *La 138.91	72 Hf 178.49	73 Ta 180.95	74 W 183.85	75 Re 186.21	76 Os 190.2	77 Ir 192.2	78 Pt 195.08	79 Au 196.97	80 Hg 200.59	81 Tl 204.38	82 Pb 207.2	83 Bi 208.98	84 Po (209)	85 At (210)	86 Rn (222)
87 Fr (223)	88 Ra 226.02	89 †Ac 227.03	104 Rf (261)	105 Db (262)	106 Sg (266)	107 Bh (264)	108 Hs (277)	109 Mt (268)	110 Ds (271)	111 Rg (272)							

*Lanthanide Series

58 Ce 140.12	59 Pr 140.91	60 Nd 144.24	61 Pm (145)	62 Sm 150.4	63 Eu 151.97	64 Gd 157.25	65 Tb 158.93	66 Dy 162.50	67 Ho 164.93	68 Er 167.26	69 Tm 168.93	70 Yb 173.04	71 Lu 174.97
90 Th 232.04	91 Pa 231.04	92 U 238.03	93 Np (237)	94 Pu (244)	95 Am (243)	96 Cm (247)	97 Bk (247)	98 Cf (251)	99 Es (252)	100 Fm (257)	101 Md (258)	102 No (259)	103 Lr (262)

†Actinide Series

ADVANCED PLACEMENT CHEMISTRY EQUATIONS AND CONSTANTS

Throughout the test the following symbols have the definitions specified unless otherwise noted.

L, mL = liter(s), milliliter(s)
 g = gram(s)
 nm = nanometer(s)
 atm = atmosphere(s)

mm Hg = millimeters of mercury
 J, kJ = joule(s), kilojoule(s)
 V = volt(s)
 mol = mole(s)

ATOMIC STRUCTURE

$$E = h\nu$$

$$c = \lambda\nu$$

E = energy
 ν = frequency
 λ = wavelength

Planck's constant, $h = 6.626 \times 10^{-34}$ J s

Speed of light, $c = 2.998 \times 10^8$ m s⁻¹

Avogadro's number = 6.022×10^{23} mol⁻¹

Electron charge, $e = -1.602 \times 10^{-19}$ coulomb

EQUILIBRIUM

$$K_c = \frac{[C]^c[D]^d}{[A]^a[B]^b}, \text{ where } aA + bB \rightleftharpoons cC + dD$$

$$K_p = \frac{(P_C)^c(P_D)^d}{(P_A)^a(P_B)^b}$$

$$K_a = \frac{[H^+][A^-]}{[HA]}$$

$$K_b = \frac{[OH^-][HB^+]}{[B]}$$

$$K_w = [H^+][OH^-] = 1.0 \times 10^{-14} \text{ at } 25^\circ\text{C}$$

$$= K_a \times K_b$$

$$\text{pH} = -\log[H^+], \text{ pOH} = -\log[OH^-]$$

$$14 = \text{pH} + \text{pOH}$$

$$\text{pH} = \text{p}K_a + \log \frac{[A^-]}{[HA]}$$

$$\text{p}K_a = -\log K_a, \text{ p}K_b = -\log K_b$$

Equilibrium Constants

K_c (molar concentrations)

K_p (gas pressures)

K_a (weak acid)

K_b (weak base)

K_w (water)

KINETICS

$$\ln[A]_t - \ln[A]_0 = -kt$$

$$\frac{1}{[A]_t} - \frac{1}{[A]_0} = kt$$

$$t_{1/2} = \frac{0.693}{k}$$

k = rate constant

t = time

$t_{1/2}$ = half-life

GASES, LIQUIDS, AND SOLUTIONS

$$PV = nRT$$

$$P_A = P_{\text{total}} \times X_A, \text{ where } X_A = \frac{\text{moles A}}{\text{total moles}}$$

$$P_{\text{total}} = P_A + P_B + P_C + \dots$$

$$n = \frac{m}{M}$$

$$K = ^\circ\text{C} + 273$$

$$D = \frac{m}{V}$$

$$KE \text{ per molecule} = \frac{1}{2}mv^2$$

Molarity, M = moles of solute per liter of solution

$$A = abc$$

P = pressure

V = volume

T = temperature

n = number of moles

m = mass

M = molar mass

D = density

KE = kinetic energy

v = velocity

A = absorbance

a = molar absorptivity

b = path length

c = concentration

Gas constant, $R = 8.314 \text{ J mol}^{-1} \text{ K}^{-1}$

$= 0.08206 \text{ L atm mol}^{-1} \text{ K}^{-1}$

$= 62.36 \text{ L torr mol}^{-1} \text{ K}^{-1}$

$1 \text{ atm} = 760 \text{ mm Hg}$

$= 760 \text{ torr}$

STP = 273.15 K and 1.0 atm

Ideal gas at STP = 22.4 L mol^{-1}

THERMOCHEMISTRY/ ELECTROCHEMISTRY

$$q = mc\Delta T$$

$$\Delta S^\circ = \sum S^\circ \text{ products} - \sum S^\circ \text{ reactants}$$

$$\Delta H^\circ = \sum \Delta H_f^\circ \text{ products} - \sum \Delta H_f^\circ \text{ reactants}$$

$$\Delta G^\circ = \sum \Delta G_f^\circ \text{ products} - \sum \Delta G_f^\circ \text{ reactants}$$

$$\Delta G^\circ = \Delta H^\circ - T\Delta S^\circ$$

$$= -RT \ln K$$

$$= -nFE^\circ$$

$$I = \frac{q}{t}$$

q = heat

m = mass

c = specific heat capacity

T = temperature

S° = standard entropy

H° = standard enthalpy

G° = standard free energy

n = number of moles

E° = standard reduction potential

I = current (amperes)

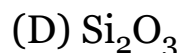
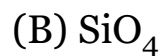
q = charge (coulombs)

t = time (seconds)

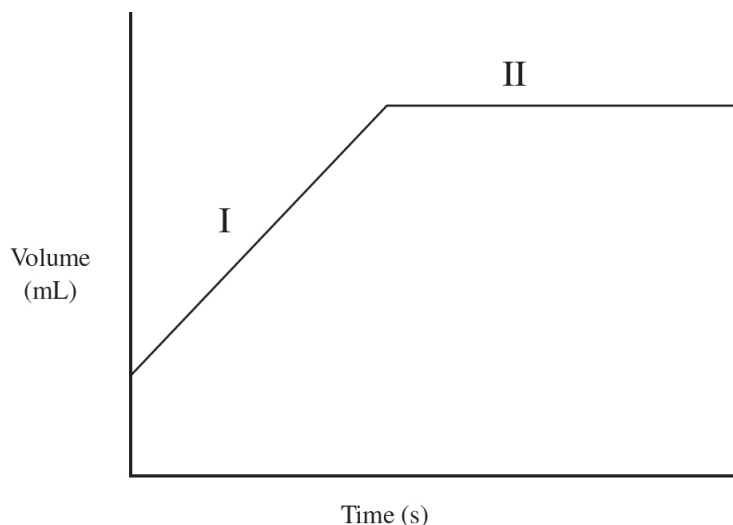
Faraday's constant, $F = 96,485 \text{ coulombs per mole of electrons}$

$$1 \text{ volt} = \frac{1 \text{ joule}}{1 \text{ coulomb}}$$

1. A compound is made up of entirely silicon and oxygen atoms. If there are 14.0 g of silicon and 32.0 g of oxygen present, what is the empirical formula of the compound?



2.



The volume of a gas is charted over time, giving the above results. Which of the following options provides a possible explanation of what was happening to the gas during each phase of the graph?

- (A) During phase I, the temperature decreased while the pressure increased. During phase II, the temperature was held constant as the pressure decreased.
- (B) During phase I, the temperature increased while the pressure was held constant. During phase II, the temperature and

pressure both decreased.

- (C) During phase I, the temperature was held constant while the pressure increased. During phase II, the temperature and pressure both decreased.
- (D) During phase I, the temperature and pressure both increased. During phase II, the temperature was held constant while the pressure decreased.

3. A solution of sulfurous acid, H_2SO_3 , is present in an aqueous solution. Which of the following represents the concentrations of three different ions in solution?

- (A) $[\text{SO}_3^{2-}] > [\text{HSO}_3^-] > [\text{H}_2\text{SO}_3]$
- (B) $[\text{H}_2\text{SO}_3] > [\text{HSO}_3^-] > [\text{SO}_3^{2-}]$
- (C) $[\text{H}_2\text{SO}_3] > [\text{HSO}_3^-] = [\text{SO}_3^{2-}]$
- (D) $[\text{SO}_3^{2-}] = [\text{HSO}_3^-] > [\text{H}_2\text{SO}_3]$

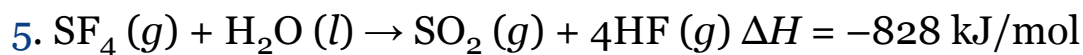


The above experiment was performed several times, and the following data was gathered:

Trial	$[\text{NO}]_{\text{init}}$ (M)	$[\text{Br}_2]_{\text{init}}$ (M)	Initial Rate of Reaction (M/min)
1	0.20 M	0.10 M	5.20×10^{-3}
2	0.20 M	0.20 M	1.04×10^{-2}
3	0.40 M	0.10 M	2.08×10^{-2}

What is the rate law for this reaction?

- (A) $\text{Rate} = k[\text{NO}][\text{Br}_2]^2$
- (B) $\text{Rate} = k[\text{NO}]^2[\text{Br}_2]^2$
- (C) $\text{Rate} = k[\text{NO}][\text{Br}_2]$
- (D) $\text{Rate} = k[\text{NO}]^2[\text{Br}_2]$



Which of the following statements accurately describes the above reaction?

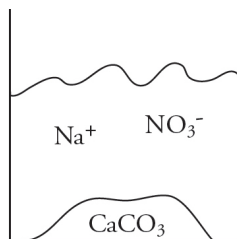
- (A) The entropy of the reactants exceeds that of the products.
- (B) $\text{H}_2\text{O}(l)$ will always be the limiting reagent.
- (C) This reaction is never thermodynamically favored.
- (D) The temperature of the surroundings will increase as this reaction progresses.

Use the following information to answer questions 6-9.

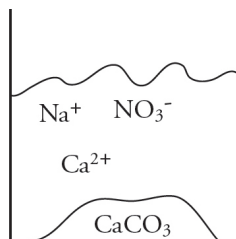
20.0 mL of 1.0 M Na_2CO_3 is placed in a beaker and titrated with a solution of 1.0 M $\text{Ca}(\text{NO}_3)_2$, resulting in the creation of a precipitate.

6. How much $\text{Ca}(\text{NO}_3)_2$ must be added to reach the equivalence point?
- (A) 10.0 mL
 - (B) 20.0 mL
 - (C) 30.0 mL
 - (D) 40.0 mL

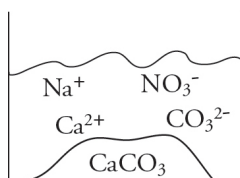
7. Which of the following diagrams correctly shows the species present in the solution in significant amounts at the equivalence point?



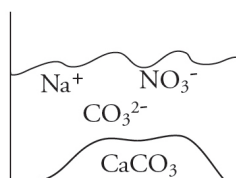
(A)



(B)



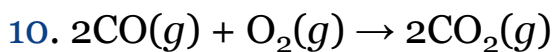
(C)



(D)

8. What will happen to the conductivity of the solution after additional $\text{Ca}(\text{NO}_3)_2$ is added past the equivalence point?
- (A) The conductivity will increase as additional ions are being added to the solution.
- (B) The conductivity will stay constant as the precipitation reaction has gone to completion.
- (C) The conductivity will decrease as the solution will be diluted with the addition of additional $\text{Ca}(\text{NO}_3)_2$.
- (D) The conductivity will stay constant as equilibrium has been established.
9. If the experiment were repeated and the Na_2CO_3 was diluted to 40.0 mL with distilled water prior to the titration, how would that affect the volume of $\text{Ca}(\text{NO}_3)_2$ needed to reach the equivalence point?
- (A) It would be cut in half.
- (B) It would decrease by a factor of 1.5.

- (C) It would double.
- (D) It would not change.



2.0 mol of CO (*g*) and 2.0 mol of O₂ (*g*) are pumped into a rigid, evacuated 4.0-L container, where they react to form CO₂ (*g*).

Which of the following values does NOT represent a potential set of concentrations for each gas at a given point during the reaction?

- | | CO | O ₂ | CO ₂ |
|-----|------|----------------|-----------------|
| (A) | 0.5 | 0.5 | 0 |
| (B) | 0 | 0.25 | 0.5 |
| (C) | 0.25 | 0.25 | 0.5 |
| (D) | 0.25 | 0.38 | 0.25 |

11. Neutral atoms of chlorine are bombarded by high-energy photons, causing the ejection of electrons from the various filled subshells. Electrons originally from which subshell would have the highest velocity after being ejected?

- (A) 1s
- (B) 2p
- (C) 3p
- (D) 3d

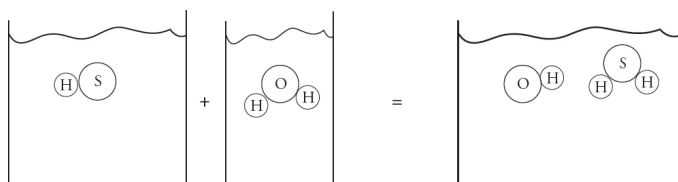
12. A sample of oxygen gas at 50°C is heated, reaching a final temperature of 100°C. Which statement best describes the behavior of the gas molecules?

- (A) Their velocity increases by a factor of two.
- (B) Their velocity increases by a factor of four.
- (C) Their kinetic energy increases by a factor of 2.
- (D) Their kinetic energy increases by a factor of less than 2.

13. The average mass, in grams, of one mole of carbon atoms is equal to

- (A) the average mass of a single carbon atom, measured in amu
- (B) the ratio of the number of carbon atoms to the mass of a single carbon atom
- (C) the number of carbon atoms in one amu of carbon
- (D) the mass, in grams, of the most abundant isotope of carbon

14.



The diagram above best represents which type of reaction?

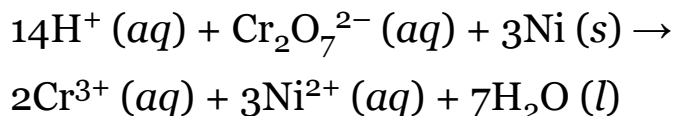
- (A) Acid/base
- (B) Oxidation/reduction
- (C) Precipitation
- (D) Decomposition

15. Which of the following is true for all bases?

- (A) All bases donate OH^- ions into solution.
- (B) Only strong bases create solutions in which OH^- ions are present.

- (C) Only strong bases are good conductors when dissolved in solution.
- (D) For weak bases, the concentration of the OH^- ions exceeds the concentration of the base in the solution.

Use the following information to answer questions 16-18.



In the above reaction, a piece of solid nickel is added to a solution of potassium dichromate.

16. Which species is being oxidized and which is being reduced?

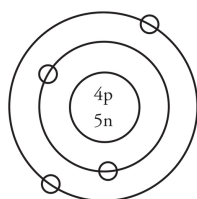
- | | Oxidized | Reduced |
|-----|-----------------------------------|-----------------------------------|
| (A) | $\text{Cr}_2\text{O}_7^{2-} (aq)$ | $\text{Ni} (s)$ |
| (B) | $\text{Cr}^{3+} (aq)$ | $\text{Ni}^{2+} (aq)$ |
| (C) | $\text{Ni} (s)$ | $\text{Cr}_2\text{O}_7^{2-} (aq)$ |
| (D) | $\text{Ni}^{2+} (aq)$ | $\text{Cr}^{3+} (aq)$ |

17. How many moles of electrons are transferred when 1 mole of potassium dichromate is mixed with 3 mol of nickel?

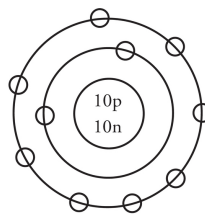
- (A) 2 moles of electrons
- (B) 3 moles of electrons
- (C) 5 moles of electrons
- (D) 6 moles of electrons

18. How does the pH of the solution change as the reaction progresses?

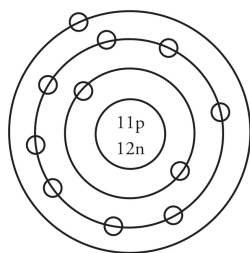
- (A) It increases until the solution becomes basic.
 (B) It increases, but the solution remains acidic.
 (C) It decreases until the solution becomes basic.
 (D) It decreases, but the solution remains acidic.
19. A sample of an unknown chloride compound was dissolved in water, and then titrated with excess $\text{Pb}(\text{NO}_3)_2$ to create a precipitate. After drying, it is determined there are 0.0050 mol of precipitate present. What mass of chloride is present in the original sample?
- (A) 0.177 g
 (B) 0.355 g
 (C) 0.522 g
 (D) 0.710 g
20. A photoelectron spectra for which of the following atoms would show peaks at exactly three different binding energies?



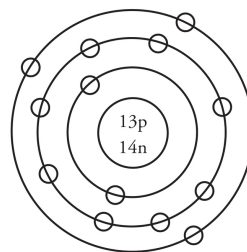
(A)



(B)



(C)



(D)

21. The bond length between any two nonmetal atoms is achieved under which of the following conditions?

- (A) Where the energy of interaction between the atoms is at its minimum value
- (B) Where the nuclei of each atom exhibits the strongest attraction to the electrons of the other atom
- (C) The point at which the attractive and repulsive forces between the two atoms are equal
- (D) The closest point at which a valence electron from one atom can transfer to the other atom

22. Hydrogen fluoride, HF, is a liquid at 15°C. All other hydrogen halides (represented by HX, where X is any other halogen) are gases at the same temperature. Why?

- (A) Fluorine has a very high electronegativity; therefore, the H-F bond is stronger than any other H-X bond.
- (B) HF is smaller than any other H-X molecule; therefore, it exhibits stronger London dispersion forces.
- (C) The dipoles in a HF molecule exhibit a particularly strong attraction force to the dipoles in other HF molecules.
- (D) The H-F bond is the most ionic in character compared to all other hydrogen halides.

23.

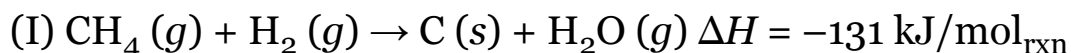
	Initial pH	pH after NaOH addition
Acid 1	3.0	3.5
Acid 2	3.0	5.0

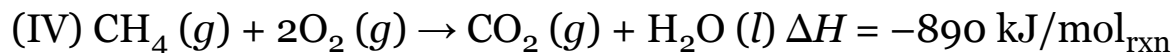
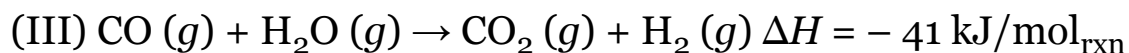
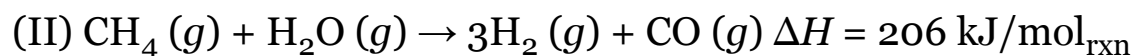
Two different acids with identical pH are placed in separate beakers. Identical portions of NaOH are added to each beaker, and the resulting pH is indicated in the table above. What can be determined about the strength of each acid?

- (A) Acid 1 is a strong acid and acid 2 is a weak acid because acid 1 resists change in pH more effectively.
 - (B) Acid 1 is a strong acid and acid 2 is a weak acid because the NaOH is more effective at neutralizing acid 2.
 - (C) Acid 1 is a weak acid and acid 2 is a strong acid because the concentration of the weak acid must be significantly greater to have the same pH as the strong acid.
 - (D) Acid 1 is a weak acid and acid 2 is a strong acid because the concentration of the hydrogen ions will be greater in acid 2 after the NaOH addition.
24. A stock solution of 12.0 *M* sulfuric acid is made available. What is the best procedure to make up 100.0 mL of 4.0 *M* sulfuric acid using the stock solution and water prior to mixing?
- (A) Add 33.3 mL of water to the flask, and then add 66.7 mL of 12.0 *M* acid.
 - (B) Add 33.3 mL of 12.0 *M* acid to the flask, and then dilute it with 66.7 mL of water.
 - (C) Add 67.7 mL of 12.0 *M* acid to the flask, and then dilute it with 33.3 mL of water.
 - (D) Add 67.7 mL of water to the flask, and then add 33.3 mL of 12.0 *M* acid.

Use the following data to answer questions 25-29.

The enthalpy values for several reactions are as follows:





25. In which of the reactions does the amount of energy released by the formation of bonds in the products exceed the amount of energy necessary to break the bonds of the reactants by the greatest amount?

- (A) Reaction I
- (B) Reaction II
- (C) Reaction III
- (D) Reaction IV

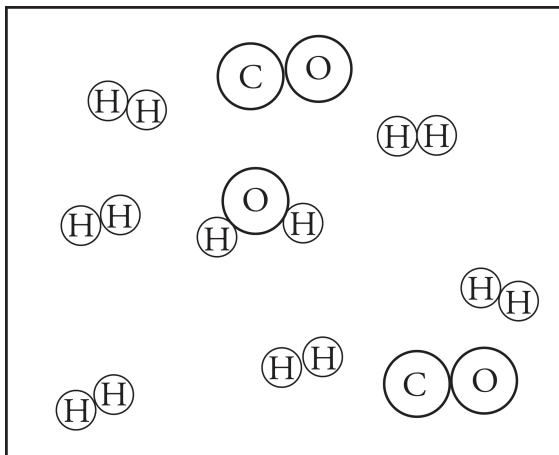
26. In which of the reactions is the value for ΔS the most positive?

- (A) Reaction I
- (B) Reaction II
- (C) Reaction III
- (D) Reaction IV

27. Regarding reaction I, how would the addition of a catalyst affect the enthalpy and entropy changes for this reaction?

- | | Enthalpy | Entropy |
|-----|-----------|-----------|
| (A) | Decrease | Decrease |
| (B) | Decrease | No Change |
| (C) | No Change | Decrease |
| (D) | No Change | No Change |

28.



Regarding reaction II, to achieve the products present in the above diagram how many moles of each reactant must be present prior to the reaction?

- (A) 1.0 mol of CH_4 and 2.0 mol of H_2O
- (B) 2.0 mol of CH_4 and 2.0 mol of H_2O
- (C) 2.0 mol of CH_4 and 3.0 mol of H_2O
- (D) 3.0 mol of CH_4 and 2.0 mol of H_2O

29. Regarding reaction IV, how much heat is absorbed or released when 2.0 mol of $\text{CH}_4(g)$ reacts with 2.0 mol of $\text{O}_2(g)$?

- (A) 890 kJ of heat is released.
- (B) 890 kJ of heat is absorbed.
- (C) 1780 kJ of heat is released.
- (D) 1780 kJ of heat is absorbed.

30. London dispersion forces are caused by

- (A) temporary dipoles created by the position of electrons around the nuclei in a molecule

- (B) the three-dimensional intermolecular bonding present in all covalent substances
- (C) the uneven electron-to-proton ratio found on individual atoms of a molecule
- (D) the electronegativity differences between the different atoms in a molecule

31. What is the general relationship between temperature and entropy for diatomic gases?

- (A) They are completely independent of each other; temperature has no effect on entropy.
- (B) There is a direct relationship, because at higher temperatures there is an increase in energy dispersal.
- (C) There is an inverse relationship, because at higher temperatures substances are more likely to be in a gaseous state.
- (D) It depends on the specific gas and the strength of the intermolecular forces between individual molecules.

32. Mixing equimolar amounts of which of the following solutions would create a buffer with a pH between 4 and 5?

$$K_a \text{ for HC}_3\text{H}_2\text{O}_2 = 1.75 \times 10^{-5}$$

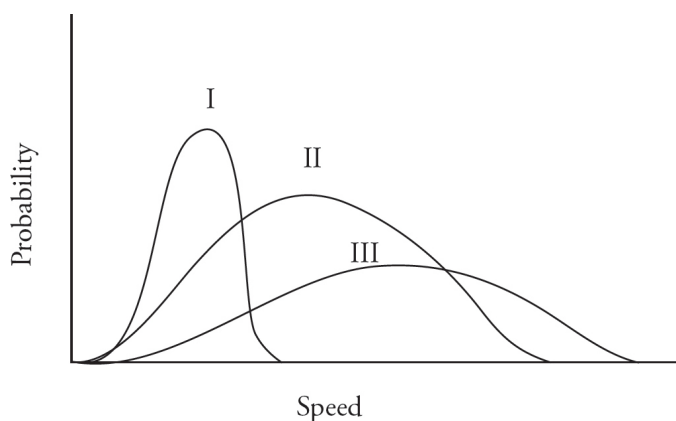
$$K_a \text{ for HPO}_4^{2-} = 4.8 \times 10^{-13}$$

- (A) H_2SO_4 and H_2PO_4
- (B) HPO_4^{2-} and Na_3PO_4
- (C) $\text{HC}_3\text{H}_2\text{O}_2$ and $\text{NaC}_3\text{H}_2\text{O}_2$
- (D) NaOH and $\text{HC}_2\text{H}_3\text{O}_2$

33. A solution contains a mixture of four different compounds: KCl (aq), $\text{Fe}(\text{NO}_3)_3$ (aq), MgSO_4 (aq), and N_2H_4 (aq). Which of these compounds would be easiest to separate via distillation?

- (A) KCl (aq)
- (B) $\text{Fe}(\text{NO}_3)_3$ (aq)
- (C) MgSO_4 (aq)
- (D) N_2H_4 (aq)

34.



Identify the three gases represented on the Maxwell-Boltzmann diagram above. Assume all gases are at the same temperature.

- | | I | II | III |
|-----|--------------|--------------|--------------|
| (A) | H_2 | N_2 | F_2 |
| (B) | H_2 | F_2 | N_2 |
| (C) | F_2 | N_2 | H_2 |
| (D) | N_2 | F_2 | H_2 |

35. A sample of solid MgCl_2 would be most soluble in which of the following solutions?

- (A) $\text{LiOH} (aq)$
- (B) $\text{CBr}_4 (aq)$
- (C) $\text{Mg}(\text{NO}_3)_2 (aq)$
- (D) $\text{AlCl}_3 (aq)$

36. Most transition metals share a common oxidation state of +2. Which of the following best explains why?

- (A) Transition metals all have a minimum of two unpaired electrons.
- (B) Transition metals have unstable configurations and are very reactive.
- (C) Transition metals tend to gain electrons when reacting with other elements.
- (D) Transition metals will lose their outermost s-block electrons when forming bonds.

37. $2\text{Ag}^+ (aq) + \text{Fe} (s) \rightarrow 2\text{Ag} (s) + \text{Fe}^{2+} (aq)$

Which of the following would cause an increase in potential in the voltaic cell described by the above reaction?

- (A) Increasing $[\text{Fe}^{2+}]$
- (B) Adding more $\text{Fe} (s)$
- (C) Decreasing $[\text{Fe}^{2+}]$
- (D) Removing some $\text{Fe} (s)$

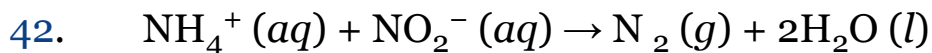
Use the following information to answer questions 38-41.

Consider the Lewis structures for the following molecules:



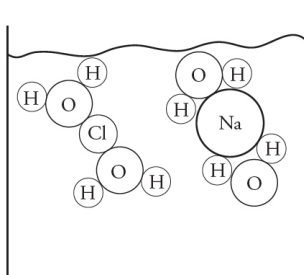
38. Which molecule would have the shortest bonds?
- (A) CO_2
 - (B) CO_3^{2-}
 - (C) NO_2^-
 - (D) NO_3^-
39. Which molecules are best represented by multiple resonance structures?
- (A) CO_2 and CO_3^{2-}
 - (B) NO_2^- and NO_3^-
 - (C) CO_3^{2-} and NO_3^-
 - (D) CO_3^{2-} , NO_2^- , and NO_3^-
40. Which molecule or molecules exhibit sp^2 hybridization around the central atom?
- (A) CO_2 and CO_3^{2-}
 - (B) NO_2^- and NO_3^-
 - (C) CO_3^{2-} and NO_3^-
 - (D) CO_3^{2-} , NO_2^- , and NO_3^-
41. Which molecule would have the smallest bond angle between terminal atoms?

- (A) CO_2
- (B) CO_3^{2-}
- (C) NO_2^-
- (D) NO_3^-

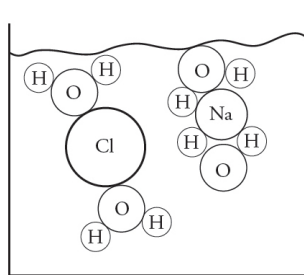


Increasing the temperature of the above reaction will increase the rate of reaction. Which of the following is NOT a reason that increased temperature increases reaction rate?

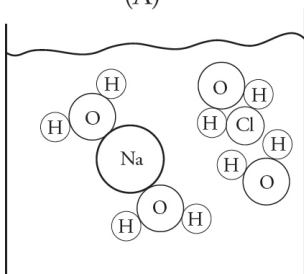
- (A) The reactants will be more likely to overcome the activation energy.
 - (B) The number of collisions between reactant molecules will increase.
 - (C) A greater distribution of reactant molecules will have high velocities.
 - (D) Alternate reaction pathways become available at higher temperatures.
43. Which of the following diagrams best represents what is happening on a molecular level when NaCl dissolves in water?



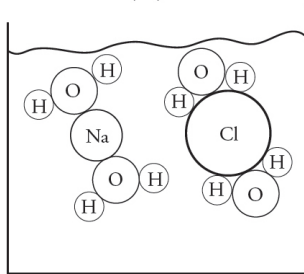
(A)



(C)



(B)



(D)

44. Nitrous acid, HNO_2 , has a $\text{p}K_a$ value of 3.3. If a solution of nitrous acid is found to have a pH of 4.2, what can be said about the concentration of the conjugate acid/base pair found in solution?

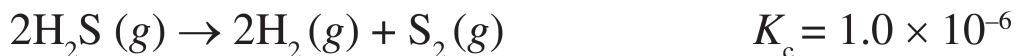
- (A) $[\text{HNO}_2] > [\text{NO}_2^-]$
- (B) $[\text{NO}_2^-] > [\text{HNO}_2]$
- (C) $[\text{H}_2\text{NO}_2^+] > [\text{HNO}_2]$
- (D) $[\text{HNO}_2] > [\text{H}_2\text{NO}_2^+]$

45. Which of the following processes is an irreversible reaction?

- (A) $\text{CH}_4 (g) + \text{O}_2 (g) \rightarrow \text{CO}_2 (g) + \text{H}_2\text{O} (l)$
- (B) $\text{HCN} (aq) + \text{H}_2\text{O} (l) \rightarrow \text{CN}^- (aq) + \text{H}_3\text{O}^+ (aq)$
- (C) $\text{Al}(\text{NO}_3)_3 (s) \rightarrow \text{Al}^{3+} (aq) + 3\text{NO}_3^- (aq)$
- (D) $2\text{Ag}^+ (aq) + \text{Ti} (s) \rightarrow 2\text{Ag} (s) + \text{Ti}^{2+} (aq)$

Use the following information to answer questions 46-50.

A sample of H_2S gas is placed in an evacuated, sealed container and heated until the following decomposition reaction occurs at 1000 K:



46. Which of the following represents the equilibrium constant for this reaction?

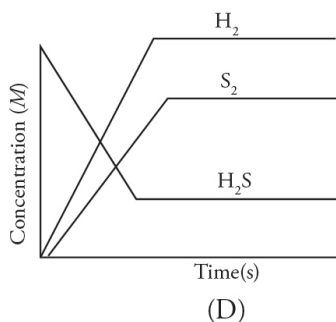
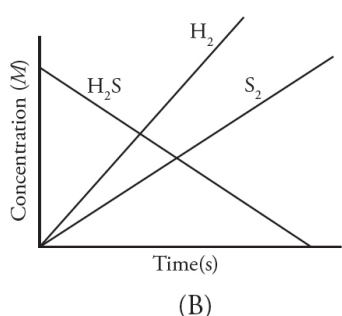
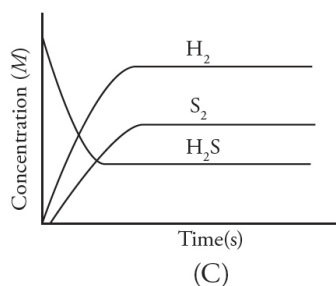
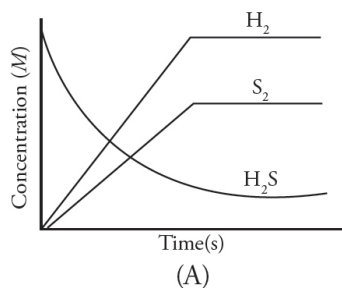
(A) $K_c = \frac{[\text{H}_2]^2 [\text{S}_2]}{[\text{H}_2\text{S}]^2}$

(B) $K_c = \frac{[\text{H}_2\text{S}]^2}{[\text{H}_2]^2 [\text{S}_2]}$

(C) $K_c = \frac{2[\text{H}_2][\text{S}_2]}{2[\text{H}_2\text{S}]}$

(D) $K_c = \frac{2[\text{H}_2\text{S}]}{2[\text{H}_2][\text{S}_2]}$

47. Which of the following graphs would best represent the change in concentration of the various species involved in the reaction over time?

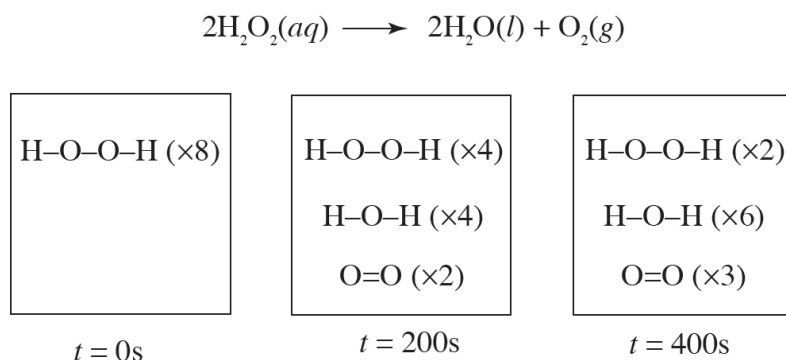


48. Which option best describes what will immediately occur to the reaction rates if the pressure on the system is increased after it has reached equilibrium?
- (A) The rate of both the forward and reverse reactions will increase.
 - (B) The rate of the forward reaction will increase while the rate of the reverse reaction decreases.
 - (C) The rate of the forward reaction will decrease while the rate of the reverse reaction increases.
 - (D) Neither the rate of the forward nor reverse reactions will change.
49. If, at a given point in the reaction, the value for the reaction quotient Q is determined to be 2.5×10^{-8} , which of the following is occurring?
- (A) The concentration of the reactant is decreasing while the concentration of the products is increasing.

- (B) The concentration of the reactant is increasing while the concentration of the products is decreasing.
 - (C) The system has passed the equilibrium point, and the concentration of all species involved in the reaction will remain constant.
 - (D) The concentrations of all species involved are changing at the same rate.
50. As the reaction progresses at a constant temperature of 1000 K, how does the value for the Gibbs free energy constant for the reaction change?
- (A) It stays constant.
 - (B) It increases exponentially.
 - (C) It increases linearly.
 - (D) It decreases exponentially.
51. An unknown substance is found to have a high melting point. In addition, it is a poor conductor of electricity and does not dissolve in water. The substance most likely contains
- (A) ionic bonding
 - (B) nonpolar covalent bonding
 - (C) covalent network bonding
 - (D) metallic bonding
52. Which of the following best explains why the ionization of atoms can occur during photoelectron spectroscopy, even though ionization is not a thermodynamically favored process?
- (A) It is an exothermic process due to the release of energy as an electron is liberated from the Coulombic attraction holding it to the nucleus.

- (B) The entropy of the system increases due to the separation of the electron from its atom.
- (C) Energy contained in the light can be used to overcome the Coulombic attraction between electrons and the nucleus.
- (D) The products of the ionization are at a lower energy state than the reactants.

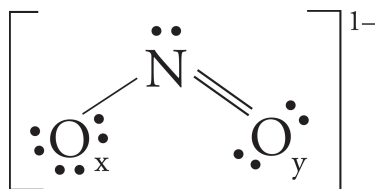
53.



The above diagrams show the decomposition of hydrogen peroxide in a sealed container in the presence of a catalyst. What is the overall order for the reaction?

- (A) Zero order
- (B) First order
- (C) Second order
- (D) Third order

54.



One of the resonance structures for the nitrite ion is shown above. What is the formal charge on each atom?

O_x N O_y

(A) -1 $+1$ -1

(B) $+1$ -1 0

(C) 0 0 -1

(D) -1 0 0

Use the following information to answer questions 55-57.

Atoms of four elements are examined: carbon, nitrogen, neon, and sulfur.

55. Atoms of which element would have the strongest magnetic moment?

(A) Carbon

(B) Nitrogen

(C) Neon

(D) Sulfur

56. Atoms of which element are most likely to form a structure with the formula XF_6 (where X is one of the four atoms)?

(A) Carbon

(B) Nitrogen

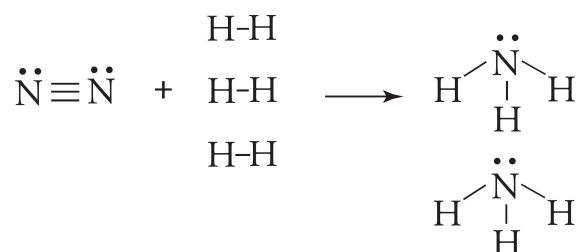
(C) Neon

(D) Sulfur

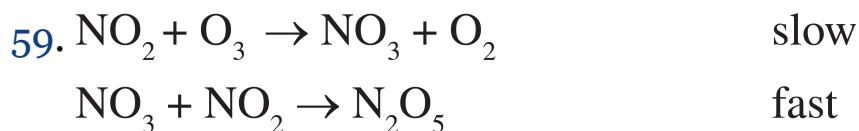
57. Which element would have a photoelectron spectra in which the peak representing electrons with the lowest ionization energy would be three times higher than all other peaks?

- (A) Carbon
- (B) Nitrogen
- (C) Neon
- (D) Sulfur

58. The diagram below supports which of the following conclusions about the reaction shown below?



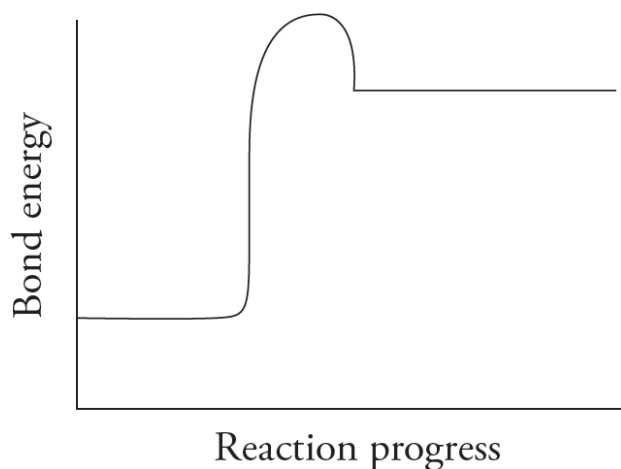
- (A) There is an increase in entropy.
- (B) Mass is conserved in all chemical reactions.
- (C) The pressure increases after the reaction goes to completion.
- (D) The enthalpy value is positive.



A proposed reaction mechanism for the reaction of nitrogen dioxide and ozone is detailed above. Which of the following is the rate law for the reaction?

- (A) $\text{Rate} = k[\text{NO}_2][\text{O}_3]$
- (B) $\text{Rate} = k[\text{NO}_3][\text{NO}_2]$
- (C) $\text{Rate} = k[\text{NO}_2]^2[\text{O}_3]$
- (D) $\text{Rate} = k[\text{NO}_3][\text{O}_2]$

60.



The concentrations of the reactants and products in the reaction represented by the above graph are found to be changing very slowly. Which of the following statements best describes the reaction given that the reaction is exergonic? ($\Delta G < 0$)

- (A) The reaction is under kinetic control.
- (B) The reaction has reached a state of equilibrium.
- (C) The reaction is highly exothermic in nature.
- (D) The addition of heat will increase the rate of reaction significantly.

END OF SECTION I

Section II

INFORMATION IN THE TABLE BELOW AND ON THE FOLLOWING PAGES MAY BE USEFUL IN ANSWERING THE QUESTIONS IN THIS SECTION OF THE EXAMINATION.

DO NOT DETACH FROM BOOK.

PERIODIC TABLE OF THE ELEMENTS

1 H 1.008																	2 He 4.00
3 Li 6.94	4 Be 9.01											5 B 10.81	6 C 12.01	7 N 14.01	8 O 16.00	9 F 19.00	10 Ne 20.18
11 Na 22.99	12 Mg 24.30											13 Al 26.98	14 Si 28.09	15 P 30.97	16 S 32.06	17 Cl 35.45	18 Ar 39.95
19 K 39.10	20 Ca 40.08	21 Sc 44.96	22 Ti 47.90	23 V 50.94	24 Cr 52.00	25 Mn 54.94	26 Fe 55.85	27 Co 58.93	28 Ni 58.69	29 Cu 63.55	30 Zn 65.39	31 Ga 69.72	32 Ge 72.59	33 As 74.92	34 Se 78.96	35 Br 79.90	36 Kr 83.80
37 Rb 85.47	38 Sr 87.62	39 Y 88.91	40 Zr 91.22	41 Nb 92.91	42 Mo 95.94	43 Tc (98)	44 Ru 101.1	45 Rh 102.91	46 Pd 106.42	47 Ag 107.87	48 Cd 112.41	49 In 114.82	50 Sn 118.71	51 Sb 121.75	52 Te 127.60	53 I 126.91	54 Xe 131.29
55 Cs 132.91	56 Ba 137.33	57 *La 138.91	72 Hf 178.49	73 Ta 180.95	74 W 183.85	75 Re 186.21	76 Os 190.2	77 Ir 192.2	78 Pt 195.08	79 Au 196.97	80 Hg 200.59	81 Tl 204.38	82 Pb 207.2	83 Bi 208.98	84 Po (209)	85 At (210)	86 Rn (222)
87 Fr (223)	88 Ra 226.02	89 †Ac 227.03	104 Rf (261)	105 Db (262)	106 Sg (266)	107 Bh (264)	108 Hs (277)	109 Mt (268)	110 Ds (271)	111 Rg (272)							

*Lanthanide Series	58 Ce 140.12	59 Pr 140.91	60 Nd 144.24	61 Pm (145)	62 Sm 150.4	63 Eu 151.97	64 Gd 157.25	65 Tb 158.93	66 Dy 162.50	67 Ho 164.93	68 Er 167.26	69 Tm 168.93	70 Yb 173.04	71 Lu 174.97
†Actinide Series	90 Th 232.04	91 Pa 231.04	92 U 238.03	93 Np (237)	94 Pu (244)	95 Am (243)	96 Cm (247)	97 Bk (247)	98 Cf (251)	99 Es (252)	100 Fm (257)	101 Md (258)	102 No (259)	103 Lr (262)

ADVANCED PLACEMENT CHEMISTRY EQUATIONS AND CONSTANTS

Throughout the test the following symbols have the definitions specified unless otherwise noted.

L, mL = liter(s), milliliter(s)
 g = gram(s)
 nm = nanometer(s)
 atm = atmosphere(s)

mm Hg = millimeters of mercury
 J, kJ = joule(s), kilojoule(s)
 V = volt(s)
 mol = mole(s)

ATOMIC STRUCTURE

$$E = h\nu$$

$$c = \lambda\nu$$

E = energy
 ν = frequency
 λ = wavelength

Planck's constant, $h = 6.626 \times 10^{-34}$ J s

Speed of light, $c = 2.998 \times 10^8$ m s⁻¹

Avogadro's number = 6.022×10^{23} mol⁻¹

Electron charge, $e = -1.602 \times 10^{-19}$ coulomb

EQUILIBRIUM

$$K_c = \frac{[C]^c[D]^d}{[A]^a[B]^b}, \text{ where } aA + bB \rightleftharpoons cC + dD$$

$$K_p = \frac{(P_C)^c(P_D)^d}{(P_A)^a(P_B)^b}$$

$$K_a = \frac{[H^+][A^-]}{[HA]}$$

$$K_b = \frac{[OH^-][HB^+]}{[B]}$$

$$K_w = [H^+][OH^-] = 1.0 \times 10^{-14} \text{ at } 25^\circ\text{C}$$

$$= K_a \times K_b$$

$$\text{pH} = -\log[H^+], \text{ pOH} = -\log[OH^-]$$

$$14 = \text{pH} + \text{pOH}$$

$$\text{pH} = \text{p}K_a + \log \frac{[A^-]}{[HA]}$$

$$\text{p}K_a = -\log K_a, \text{ p}K_b = -\log K_b$$

Equilibrium Constants

K_c (molar concentrations)

K_p (gas pressures)

K_a (weak acid)

K_b (weak base)

K_w (water)

KINETICS

$$\ln[A]_t - \ln[A]_0 = -kt$$

$$\frac{1}{[A]_t} - \frac{1}{[A]_0} = kt$$

$$t_{1/2} = \frac{0.693}{k}$$

k = rate constant

t = time

$t_{1/2}$ = half-life

GASES, LIQUIDS, AND SOLUTIONS

$$PV = nRT$$

$$P_A = P_{\text{total}} \times X_A, \text{ where } X_A = \frac{\text{moles A}}{\text{total moles}}$$

$$P_{\text{total}} = P_A + P_B + P_C + \dots$$

$$n = \frac{m}{M}$$

$$K = ^\circ\text{C} + 273$$

$$D = \frac{m}{V}$$

$$KE \text{ per molecule} = \frac{1}{2}mv^2$$

Molarity, M = moles of solute per liter of solution

$$A = abc$$

P = pressure

V = volume

T = temperature

n = number of moles

m = mass

M = molar mass

D = density

KE = kinetic energy

v = velocity

A = absorbance

a = molar absorptivity

b = path length

c = concentration

Gas constant, $R = 8.314 \text{ J mol}^{-1} \text{ K}^{-1}$

$= 0.08206 \text{ L atm mol}^{-1} \text{ K}^{-1}$

$= 62.36 \text{ L torr mol}^{-1} \text{ K}^{-1}$

$1 \text{ atm} = 760 \text{ mm Hg}$

$= 760 \text{ torr}$

STP = 273.15 K and 1.0 atm

Ideal gas at STP = 22.4 L mol^{-1}

THERMOCHEMISTRY/ ELECTROCHEMISTRY

$$q = mc\Delta T$$

$$\Delta S^\circ = \sum S^\circ \text{ products} - \sum S^\circ \text{ reactants}$$

$$\Delta H^\circ = \sum \Delta H_f^\circ \text{ products} - \sum \Delta H_f^\circ \text{ reactants}$$

$$\Delta G^\circ = \sum \Delta G_f^\circ \text{ products} - \sum \Delta G_f^\circ \text{ reactants}$$

$$\Delta G^\circ = \Delta H^\circ - T\Delta S^\circ$$

$$= -RT \ln K$$

$$= -nFE^\circ$$

$$I = \frac{q}{t}$$

q = heat

m = mass

c = specific heat capacity

T = temperature

S° = standard entropy

H° = standard enthalpy

G° = standard free energy

n = number of moles

E° = standard reduction potential

I = current (amperes)

q = charge (coulombs)

t = time (seconds)

Faraday's constant, $F = 96,485 \text{ coulombs per mole of electrons}$

$$1 \text{ volt} = \frac{1 \text{ joule}}{1 \text{ coulomb}}$$

CHEMISTRY
SECTION II
7 Questions

(Total time—1 hour and 45 minutes)

YOU MAY USE YOUR CALCULATOR FOR THIS SECTION.

Directions: Questions 1-3 are long free-response questions that require about 23 minutes each to answer and are worth 10 points each. Questions 4-7 are short free-response questions that require about 9 minutes each to answer and are worth 4 points each.

Write your response in the space provided following each question. Examples and equations may be included in your responses where appropriate. For calculations, clearly show the method used and the steps involved in arriving at your answers. You must show your work to receive credit for your answer. Pay attention to significant figures.

1. A stock solution of 0.100 *M* cobalt (II) chloride is used to create several solutions, indicated in the data table below:

Sample	Volume CoCl₂ (mL)	Volume H₂O (mL)
1	20.00	0
2	15.00	5.00
3	10.00	10.00
4	5.00	15.00

- (a) In order to achieve the degree of accuracy shown in the table above, select which of the following pieces of laboratory equipment could be used when measuring out the CoCl₂:

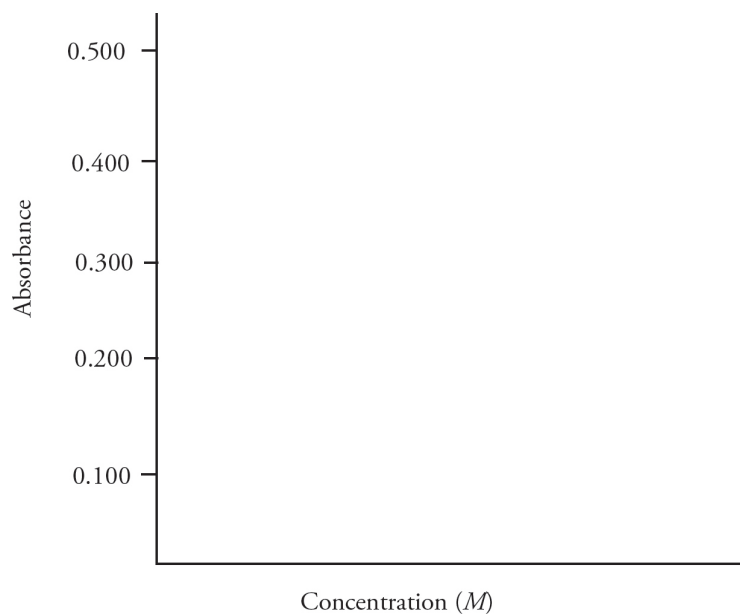
- 150-mL beaker
- 400-mL beaker
- 250-mL Erlenmeyer flask
- 50-mL buret
- 50-mL graduated cylinder
- 100-mL graduated cylinder

(b) Calculate the concentration of the CoCl_2 in each sample.

The solutions are then placed in cuvettes before being inserted into a spectrophotometer calibrated to 560 nm and their values are measured, yielding the data below:

Sample	Absorbance
1	0.485
2	0.364
3	0.243
4	0.121

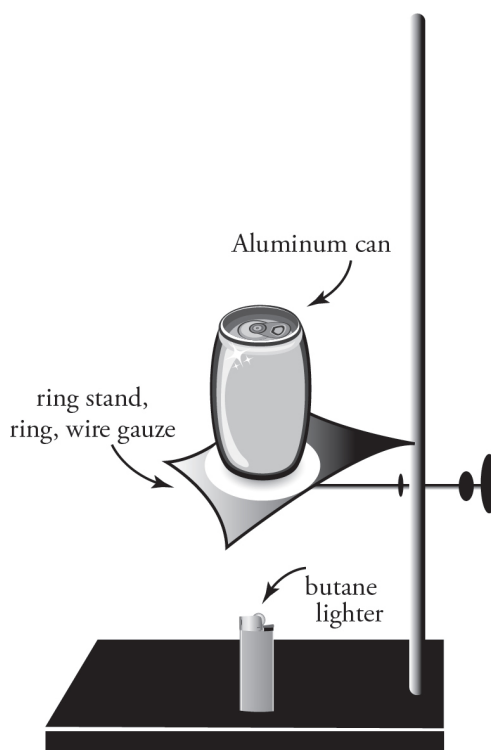
- (c) If gloves are not worn when handling the cuvettes, how might this affect the absorbance values gathered?
- (d) If the path length of the cuvette is 1.00 cm, what is the molar absorptivity value for CoCl_2 at 560 nm?
- (e) On the axes on the next page, plot a graph of absorbance vs. concentration. The y -axes scale is set, and be sure to scale the x -axes appropriately



(f) What would the absorbance values be for CoCl_2 solutions at the following concentrations?

(i) 0.067 M

(ii) 0.180 M



2. A sample of liquid butane (C_4H_{10}) in a pressurized lighter is set up directly beneath an aluminum can, as show in the diagram above. The can contains 100.0 mL of water, and when the butane is ignited the temperature of the water inside the can increases from 25.0°C to 82.3°C . The total mass of butane ignited is found to be 0.51 g, the specific heat of water is $4.18 \text{ J/g}\cdot^\circ\text{C}$, and the density of water is 1.00 g/mL .

(a) Write the balanced chemical equation for the combustion of one mole of butane in air.

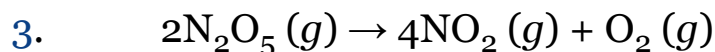
(b) (i) How much heat did the water gain?

(ii) What is the experimentally determined heat of combustion for butane based on this experiment? Your answer should be in kJ/mol .

(c) Given butane's density of 0.573 g/mL at 25°C , calculate how much heat would be emitted if 5.00 mL of it were combusted at that temperature.

(d) The overall combustion of butane is an exothermic reaction. Explain why this is, in terms of bond energies.

(e) One of the major sources of error in this experiment comes from the heat that is absorbed by the air. Why, then, might it not be a good idea to perform this experiment inside a sealed container to prevent the heat from leaving the system?



The data below was gathered for the decomposition of N_2O_5 at 310 K via the equation above.

Time (s)	$[\text{N}_2\text{O}_5] (M)$

0	0.250
500.	0.190
1000.	0.145
2000.	0.085

- (a) How does the rate of appearance of NO_2 compare to the rate of disappearance of N_2O_5 ? Justify your answer.
- (b) The reaction is determined to be first order overall. On the axes below, create a graph of some function of concentration vs. time that will produce a straight line. Label and scale your axes appropriately.



- (c) (i) What is the rate constant for this reaction? Include units.
- (ii) What would the concentration of N_2O_5 be at $t = 1500$ s?
- (iii) What is the half-life of N_2O_5 ?

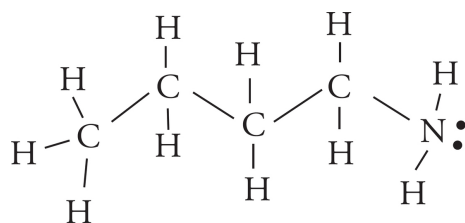
(d) Would the addition of a catalyst increase, decrease, or have no effect on the following variables? Justify your answers.

(i) Rate of disappearance of N_2O_5

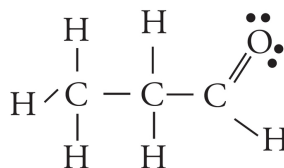
(ii) Magnitude of the rate constant

(iii) Half-life of N_2O_5

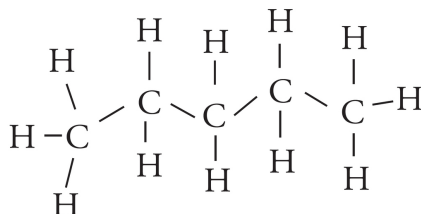
4. Consider the Lewis structures for the following four molecules:



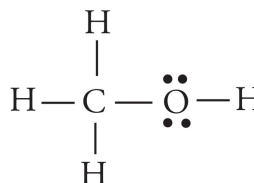
n-Butylamine



Propanal

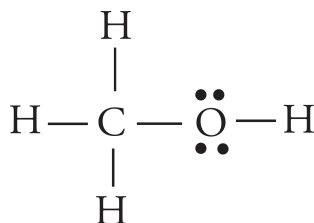


Pentane



Methanol

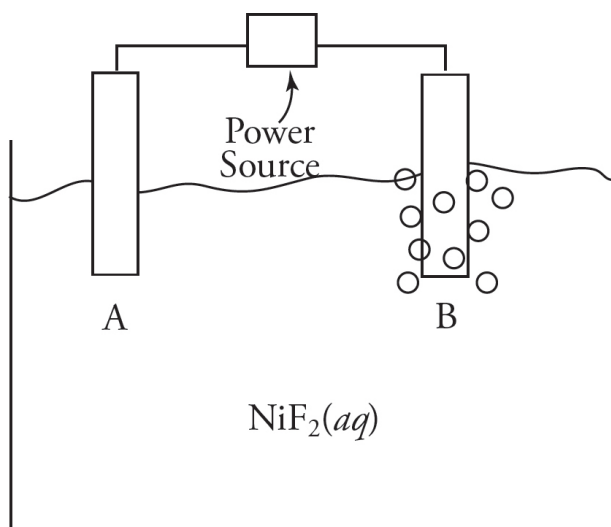
- (a) All of the substances are liquids at room temperature. Organize them from high to low in terms of boiling points, clearly differentiating between the intermolecular forces in each substance.
- (b) On the methanol diagram reproduced below, draw the locations of all dipoles.



Methanol

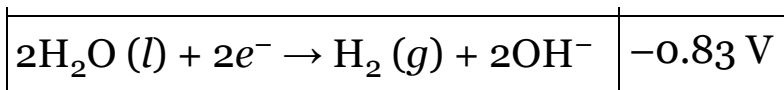
(c) n-Butylamine is found to have the lowest vapor pressure at room temperature out of the four liquids. Justify this observation in terms of intermolecular forces.

5. Current is run through an aqueous solution of nickel (II) fluoride, and a gas is evolved at the right-hand electrode, as indicated by the diagram below:



The standard reduction potential for several reactions is given in the following table:

Half-cell	E°_{red}
$\text{F}_2(g) + 2e^- \rightarrow 2\text{F}^-$	+2.87 V
$\text{O}_2(g) + 4\text{H}^+ + 4e^- \rightarrow 2\text{H}_2\text{O}(l)$	+1.23 V
$\text{Ni}^{2+} + 2e^- \rightarrow \text{Ni}(s)$	-0.25 V



- (a) Determine which half-reaction is occurring at each electrode:
- Oxidation
 - Reduction
- (b) (i) Calculate the standard cell potential of the cell.
- (ii) Calculate the Gibbs free energy value of the cell at standard conditions.
- (c) Which electrode in the diagram (A or B) is the cathode, and which is the anode? Justify your answers.
6. Aniline, $\text{C}_6\text{H}_5\text{NH}_2$, is a weak base with $K_b = 3.8 \times 10^{-10}$.
- (a) Write out the reaction that occurs when aniline reacts with water.
- (b) (i) What is the concentration of each species at equilibrium in a solution of $0.25 \text{ M } \text{C}_6\text{H}_5\text{NH}_2$?
- (ii) What is the pH value for the solution in (i)?
7. A rigid, sealed 12.00 L container is filled with 10.00 g each of three different gases: CO_2 , NO , and NH_3 . The temperature of the gases is held constant 35.0°C . Assume ideal behavior for all gases.
- (a) (i) What is the mole fraction of each gas?
- (ii) What is the partial pressure of each gas?
- (b) Out of the three gases, molecules of which gas will have the highest velocity? Why?

- (c) Name one circumstance in which the gases might deviate from ideal behavior, and clearly explain the reason for the deviation.

STOP
END OF EXAM

Practice Test 2: Answers and Explanations

PRACTICE TEST 2: MULTIPLE-CHOICE ANSWER KEY

- | | | |
|-------|-------|-------|
| 1. B | 21. A | 41. C |
| 2. B | 22. C | 42. D |
| 3. B | 23. C | 43. D |
| 4. D | 24. D | 44. B |
| 5. D | 25. D | 45. A |
| 6. B | 26. B | 46. A |
| 7. A | 27. D | 47. C |
| 8. A | 28. C | 48. C |
| 9. D | 29. A | 49. A |
| 10. C | 30. A | 50. A |
| 11. C | 31. B | 51. C |
| 12. D | 32. C | 52. C |
| 13. A | 33. D | 53. B |
| 14. A | 34. C | 54. D |
| 15. C | 35. A | 55. B |
| 16. C | 36. D | 56. D |
| 17. D | 37. C | 57. C |
| 18. B | 38. A | 58. B |
| 19. B | 39. D | 59. A |
| 20. B | 40. D | 60. A |

SECTION I—MULTIPLE-CHOICE ANSWERS AND EXPLANATIONS

1. **B**

14.0 g of Si is 0.50 mol, and 32.0 g of oxygen is 2.0 mol. Converting that to a whole number ratio gives you 1 mol of Si for every 4 moles of O.

2. **B**

In phase I, an increased temperature means the molecules are moving faster and will spread out more, leading to an increased volume. For the volume to remain constant in phase II, either both pressure and volume have to remain constant, or they both have to increase or decrease together, as they are inversely proportional.

3. **B**

Weak acids have a very low dissociation value, and for polyprotic weak acids the second dissociation is always weaker than the first.

4. **D**

Between trial 1 and 3, the concentration of NO doubled while the concentration of Br₂ held constant, and the rate went up by a factor of four. So, the reaction is second order with respect to NO. Between trial 1 and 2, the concentration of NO was held constant while the concentration of Br₂ doubled, and the rate went up by a factor of two. Thus, the reaction is first order with respect to Br₂.

5. **D**

A negative enthalpy change indicates that energy is lost to the surroundings as the reaction occurs.

6. **B**

Equivalence is reached when the conductivity of the solution reaches its minimum value, meaning as many of the precipitate ions are out of solution as possible. As the formula of the precipitate is CaCO_3 , for every mole of sodium carbonate in the beaker, one mole of calcium nitrate will be needed. As both solutions have equal molarity, equal volumes of each would be necessary to ensure the moles are equal.

7. **A**

At the equivalence point, the moles of Ca^{2+} and CO_3^{2-} in solution will be negligible.

8. **A**

The Ca^{2+} ions will have nothing to react with, so they will just remain in solution (as will the extra NO_3^- ions being added), increasing the conductivity.

9. **D**

The equivalence point is reached when there are equal moles of each reactant present. Diluting the sodium carbonate solution will not change the number of moles of Na_2CO_3 present in solution.

10. **C**

For every two moles of CO that react, one mole of O₂ will also react. If one mole of CO reacts, half a mole of O₂ will react. At that point, the concentration of CO will be (1.0 mol/4.0 L = 0.25 M) and the concentration of the O₂ will be (1.5 mol/4.0 L = 0.38 M), rendering (C) impossible.

11. **C**

The farther away an electron is from the nucleus, the less ionization energy that is required to eject it, and as a result, the electron will have more kinetic energy after it is ejected. The 3*p* subshell is the farthest one that a neutral chlorine atom would have electrons in. Beware of (D); chlorine does not have a 3*d* subshell.

12. **D**

Temperature in Kelvins is a measure of kinetic energy. When that temperature doubles, so does the kinetic energy. However, this is not a direct correlation, so while a doubling of the temperature in Celsius would increase the amount of kinetic energy, it would not double it.

13. **A**

This is a straightforward concept to understand. The average atomic mass of an element on the periodic table measures both the average mass of a mole of atoms of that element in grams, as well as being the average mass of a single atom of that element in amu.

14. **A**

A proton is transferred from the water to the HS⁻ ion, making the reaction an acid/base reaction.

15. **C**

Strong bases dissociate to donate hydroxide ions into a solution and weak bases cause the creation of hydroxide ions by taking protons from water. However, weak bases do not protonate strongly, meaning there will not be nearly as many ions in solution for a weak base as there are for a strong base. Thus, weak bases are not good conductors.

16. **C**

The $\text{Cr}_2\text{O}_7^{2-} (aq)$ gains electrons to change the oxidation state on Cr from +6 to +3. The Ni (s) loses electrons to change the oxidation state on Ni from 0 to +2.

17. **D**

In one mole of dichromate, two moles of chromium are reduced, each requiring three moles of electrons. Three moles of nickel are being oxidized, each requiring two moles of electrons. Thus, six moles of electrons are being transferred.

18. **B**

As the reaction progresses, the H^+ ions are converting to H_2O molecules. This would raise the pH of the solution. As the reaction progresses, the solution will become less acidic, but there is no mechanism for it to become basic.

19. **B**

The precipitate is PbCl_2 , and in every mole of precipitate there are two moles of chlorine ions. Thus, there are 0.010 mol of chlorine present in the precipitate, and through conservation of mass, the same number of moles of

chlorine were present in the original sample: $0.010 \text{ mol} \times 35.5 \text{ g} = 0.355 \text{ g}$.

20. **B**

Diagram B represents an atom of neon, which has three subshells: $1s$, $2s$, and $2p$. Electrons from each subshell would have a different binding energy, yielding three peaks on a PES.

21. **A**

The bond length always corresponds to the point where the potential bond energy (a balance of the attraction and repulsion forces between the two atoms) is at its minimum value.

22. **C**

The IMFs between HF molecules are hydrogen bonds, which are very strong dipoles created due to the high electronegativity value of fluorine. No other hydrogen halide exhibits hydrogen bonding, and thus they would have weaker intermolecular forces (and lower boiling points) than HF.

23. **C**

Weak acids resist changes in pH more effectively than strong acids because so many molecules of weak acid are undissociated in solution. The base must cause those molecules to dissociate before affecting the pH significantly.

24. **D**

Using $M_1V_1 = M_2V_2$, you can calculate the amount of stock solution needed. $(12.0)V_1 = (4.0)(100.0)$, so $V_1 = 33.3$ mL. When making solutions, always add acid to water (in order to more effectively absorb the heat of the exothermic reaction).

25. **D**

When calculating enthalpy, the total energy change in the reaction is always calculated by the following:

Bonds broken (reactants) – Bonds formed (products)

The more negative this value is, the more energy that was released in the bond formation of the products compared to the amount of energy necessary to break the bonds in the reactants.

26. **B**

There are two moles of gaseous reactants, and four moles of gaseous products. That means the disorder increased, yielding a positive value for ΔS .

27. **D**

The addition of a catalyst speeds up the reaction but will not affect the enthalpy or the entropy values.

28. **C**

2.0 moles of CH_4 would react with 3.0 moles of H_2O , leaving 1.0 mole left. It would also create 6.0 moles of H_2 and 2.0 moles of CO .

29. **A**

When the listed amounts react, the O_2 will be the limiting reactant, as it is used up twice as quickly (this is determined by the coefficients in the reaction).

$$\text{So: } 2.0 \text{ mol O}_2 \times \frac{1 \text{ mol}_{\text{rxn}}}{2.0 \text{ mol O}_2} \times \frac{-890 \text{ kJ}}{1 \text{ mol}_{\text{rxn}}} = -890 \text{ kJ}$$

That gives the magnitude of the energy change, and the negative sign means heat will be released.

30. **A**

London dispersion forces are created by temporary dipoles due to the constant motion of electrons in an atom or molecule.

31. **B**

The higher the temperature of any substance, the larger the range of velocities the molecules of that substance can have, and thus the more disorder the substance can have. A Maxwell-Boltzmann diagram represents this in a graphical form.

32. **C**

When equimolar amounts of an acid and its conjugate base are mixed, the pH of the buffer which is created will be the same as the $\text{p}K_{\text{a}}$ of the weak acid. The $\text{p}K_{\text{a}}$ for $\text{HC}_2\text{H}_3\text{O}_2$ is between 4 and 5.

33. **D**

Distillation involves the boiling off of substances with different boiling points. The other three compounds are all ionic, meaning that in solution they are free ions. If one

were to boil off all of the N_2H_4 and the water, the remaining ions would all mix together to form multiple precipitates. However, by heating the solution to a boiling point greater than that of N_2H_4 and lower than that of water, the N_2H_4 can be collected in a separate flask.

34. **C**

At identical temperatures, the gases would all have identical amounts of kinetic energy. In order for that to happen, the gas with the lowest mass (H_2) would have to have the highest average velocity, and the gas with the highest mass (F_2) would have to have the lowest average velocity.

35. **A**

Like dissolves like, so ionic substances dissolve best in ionic compounds, eliminating (B). Both (C) and (D) share an ion with MgCl_2 . Via the common ion effect and Le Châtelier's principle, that would reduce the solubility of the MgCl_2 in those solutions. The best choice is (A).

36. **D**

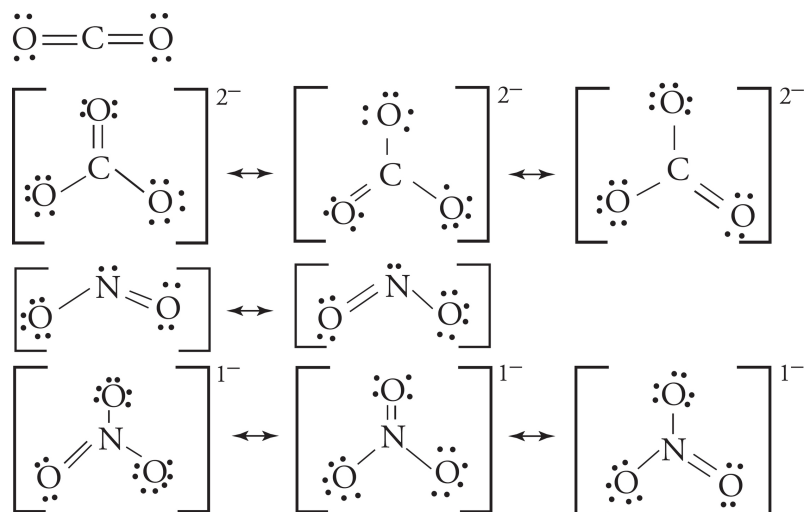
The outermost *s*-block electrons in a transition metal tend to be lost before the *d*-block electrons. Additionally, the other options do not accurately describe the properties of transition metals.

37. **C**

Adding or removing a solid would not cause any equilibrium shift. Decreasing the concentration of the Fe^{2+} causes a shift to the right, which would increase the potential of the cell.

38. **A**

The following Lewis structures are necessary to answer questions 38–41:



CO₂ has a bond order of 2, which exceeds the order of the other structures. Remember that a higher bond order corresponds with shorter and stronger bonds.

39. **D**

Except for CO₂, the other molecules all display resonance.

40. **D**

All three of those structures have three electron domains, and thus *sp*² hybridization.

41. **C**

The bond angle of NO₂[−] would be less than that of NO₃[−] or CO₃^{2−} because the unbonded pair of electrons on the nitrogen atom reduces the overall bond angle.

42. **D**

Even if this were true, any pathways that become available at higher temperatures would be less likely to be taken than the original pathway.

43. **D**

The Na^+ cations would be attracted to the negative (oxygen) end of the water molecules and the Cl^- anions would be attracted to the positive (hydrogen) end. Additionally, Na^+ ions are smaller than Cl^- ions because they have less-filled energy levels.

44. **B**

In this case, the pH is greater than the $\text{p}K_a$. This means that there will be more conjugate base present in solution than the original acid. The conjugate base of HNO_2 is NO_2^- .

45. **A**

Combustion reactions are irreversible, while the other reactions are examples of acid-base, dissolution, and oxidation-reduction, all of which are reversible in this case.

46. **A**

Equilibrium is always products over reactants, and coefficients in a balanced equilibrium reaction become exponents in the equilibrium expression.

47. **C**

The concentration of the H_2S will decrease exponentially until it reaches a constant value. The concentrations of the two products will increase exponentially (the H_2 twice as quickly as the S_2) until reaching equilibrium.

48. **C**

Increasing the pressure causes a shift toward the side with fewer gas molecules—in this case, a shift to the left. This means the reverse reaction rate increases while the forward reaction rate decreases.

49. **A**

The reaction will progress until $Q = K_c$. If $Q < K_c$, the numerator of the expression (the products) will continue to increase while the denominator (the reactant) decreases until equilibrium is established.

50. **A**

If the temperature is constant, then the equilibrium constant K is unchanged. Via $\Delta G = -RT \ln K$, if K and T are both constant, then so is the value for ΔG .

51. **C**

An ionic substance would dissolve in water, and a nonpolar covalent substance would have a low melting point. A metallic substance would be a good conductor. The only type of bonding that meets all the criteria is covalent network bonding.

52. **C**

Light contains energy (via $E = h\nu$), and that energy can be used to cause a reaction.

53. **B**

In 200 seconds, half of the original sample decayed. In another 200 seconds, half of the remaining sample

decayed. This demonstrates a first order reaction.

54. **D**

O_x has 6 valence electrons and 7 assigned electrons: $6 - 7 = -1$. Both O_y and the N atoms have the same number of valence and assigned electrons, making their formal charges zero.

55. **B**

The strength of an atom's magnetic moment increases with an increase in the number of unpaired electrons. Nitrogen has the most unpaired electrons (3), and thus the strongest magnetic moment.

56. **D**

Sulfur is the only element with an empty *d*-block in its outermost energy level, and is thus the only atom of the four that can form an expanded octet.

57. **C**

Neon has six electrons in its subshell with the lowest ionization energy (*2p*), and only two electrons in the other two subshells (*1s* and *2s*). This means the *2p* peak will be three times higher than the other two peaks.

58. **B**

The amount of matter is equal on both sides of the reaction. None of the other options are supported by the diagram.

59. **A**

The overall rate law is always equal to the rate law of the slowest elementary step. The rate law of any elementary step can be determined using the coefficients of the reactants in that step.

60. **A**

Reactions with high activation energies that do not proceed at a measurable rate are considered to be under kinetic control—that is, their rate of progress is based on kinetics instead of thermodynamics.

SECTION II—FREE-RESPONSE ANSWERS AND EXPLANATIONS

1. A stock solution of 0.100 *M* cobalt (II) chloride is used to create several solutions, indicated in the data table below:

Sample	Volume CoCl ₂ (mL)	Volume H ₂ O (mL)
1	20.00	0
2	15.00	5.00
3	10.00	10.00
4	5.00	15.00

- (a) In order to achieve the degree of accuracy shown in the table above, select which of the following pieces of laboratory equipment could be used when measuring out the CoCl₂.

- 150-mL beaker
- 400-mL beaker
- 250-mL Erlenmeyer flask
- 50-mL buret
- 50-mL graduated cylinder
- 100-mL graduated cylinder

The 50-mL buret is what is needed here in order to get measurements that are accurate to the hundredths place. Graduated cylinders are generally accurate to the tenths, and using flasks or beakers to measure out volume is highly inaccurate.

- (b) Calculate the concentration of the CoCl_2 in each sample.

The solutions are then placed in cuvettes before being inserted into a spectrophotometer calibrated to 560 nm and their values are measured, yielding the data below:

Sample	Absorbance
1	0.485
2	0.364
3	0.243
4	0.121

Sample 1: 0.100 M (no dilution occurred)

For samples 2–4, the first step is to calculate the number of moles of CoCl_2 , using the Molarity = moles/volume formula. Then, divide the number of moles by the new volume (volume of CoCl_2 + volume H_2O) to determine the new concentration.

Sample 2: 0.100 $M = n/0.015 \text{ L}$

$n = 0.0015 \text{ mol}$

$0.0015 \text{ mol}/0.020 \text{ L} = 0.075 \text{ M}$

Sample 3: 0.100 $M = n/0.010 \text{ L}$

$n = 0.0010$

$0.0010 \text{ mol}/0.020 \text{ L} = 0.050 \text{ M}$

Sample 4: 0.100 $M = n/0.0050 \text{ L}$

$n = 0.00050 \text{ mol}$

$0.00050 \text{ mol}/0.020 \text{ L} = 0.025 \text{ M}$

- (c) If gloves are not worn when handling the cuvettes, how might this affect the absorbance values gathered?

If gloves are not worn when handling the cuvettes, fingertip oils may be deposited on the surface of the cuvette. These oils may absorb some light, increasing the overall absorbance values.

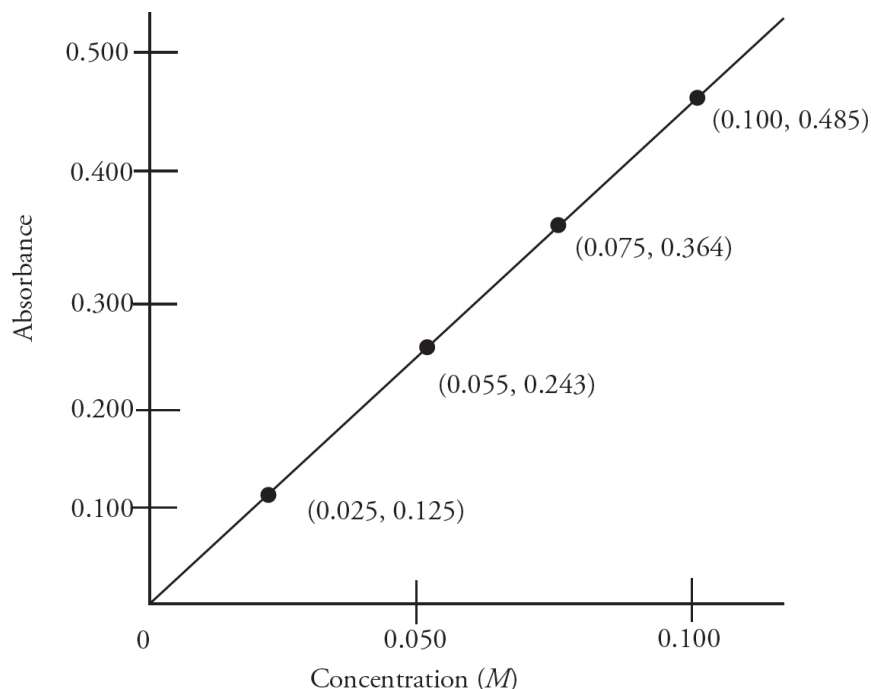
- (d) If the path length of the cuvette is 1.00 cm, what is the molar absorptivity value for CoCl_2 at 560 nm?

Using Beer's Law: $A = abc$ and the data from Sample 1:

$$0.485 = a(1.00 \text{ cm})(0.100 \text{ M}) \quad a = 4.85 \text{ M}^{-1}\text{cm}^{-1}$$

Values from any sample can be used with identical results.

- (e) On the axes below, plot a graph of absorbance vs. concentration. The y -axis scale is set, and be sure to scale the x -axis appropriately.



- (f) What would the absorbance values be for CoCl_2 solutions at the following concentrations?

While the absorbance values for (i) can be estimated from a correctly drawn graph, the absorbance value for (ii) would be off the chart. The more accurate method to determine the absorbances of both samples would be to determine the slope of the line from the graph in (e) and then plug in the appropriate values. (Note: Estimating the value for (i) is still an acceptable solution.)

Any two points can be taken to determine the slope of the graph, but by examining the Beer's Law calculation in (d) you can see it is already in slope-intercept form (the y -intercept would be zero, as at 0 concentration there would be no absorbance). So, the slope of the line is equal to $4.85 \text{ M}^{-1}\text{cm}^{-1} \times 1.00 \text{ cm} = 4.85 \text{ M}^{-1}$.

- (i) 0.067 M

$$A = (4.85 \text{ M}^{-1})(0.067 \text{ M}) = 0.325$$

- (ii) 0.180 M

$$A = (4.85 \text{ M}^{-1})(0.180 \text{ M}) = 0.873$$

2. A sample of liquid butane (C_4H_{10}) in a pressurized lighter is set up directly beneath an aluminum can, as shown in the diagram above. The can contains 100.0 mL of water, and when the butane is ignited the temperature of the water inside the can increases from 25.0°C to 82.3°C . The total mass of butane ignited is found to be 0.51 g, the specific heat of water is $4.18 \text{ J/g}\times^\circ\text{C}$, and the density of water is 1.00 g/mL .

- (a) Write the balanced chemical equation for the combustion of butane in air.

The butane reacts with the oxygen in the air, and the products of any hydrocarbon combustion are always carbon dioxide and water: $\text{C}_4\text{H}_{10}(\text{g}) + 13/2\text{O}_2(\text{g}) \rightarrow 4\text{CO}_2(\text{g}) + 5\text{H}_2\text{O}(\text{l})$.

(b) (i) How much heat did the water gain?

The formula needed is $q = mc\Delta T$.

The mass in the equation is the mass of the water, which is equal to 100.0 g (as the density of water is 1.0 g/mL). $\Delta T = 82.3^\circ\text{C} - 25.0^\circ\text{C} = 57.3^\circ\text{C}$ and c is given as 4.18 J/g $^\circ\text{C}$, so:

$$q = (100.0 \text{ g})(4.18 \text{ J/g}\cdot^\circ\text{C})(57.3^\circ\text{C}) = 24,000 \text{ J or } 24 \text{ kJ}$$

(ii) What is the experimentally determined heat of combustion for butane based on this experiment? Your answer should be in kJ/mol.

$$0.51 \text{ g C}_4\text{H}_{10} \times 1 \text{ mol C}_4\text{H}_{10}/58.14 \text{ g C}_4\text{H}_{10} = 0.0088 \text{ mol C}_4\text{H}_{10}$$

The sign on the heat calculated in (b)(i) needs to be flipped, as the combustion reaction will generate as much heat as the water gained.

$$-24 \text{ kJ}/0.0088 \text{ mol C}_4\text{H}_{10} = -2,700 \text{ kJ/mol}$$

(c) Given butane's density of 0.573 g/mL at 25 $^\circ\text{C}$, calculate how much heat would be emitted if 5.00 mL of it were combusted at that temperature.

$$0.573 \text{ g/mL} = m/5.00 \text{ mL} \quad m = 2.87 \text{ g butane}$$

$$2.87 \text{ g C}_4\text{H}_{10} \times 1 \text{ mol C}_4\text{H}_{10}/58.14 \text{ g C}_4\text{H}_{10} = 0.049 \text{ mol C}_4\text{H}_{10}$$

$$0.049 \text{ mol C}_4\text{H}_{10} \times 2700 \text{ kJ/mol} = 130 \text{ kJ of heat emitted}$$

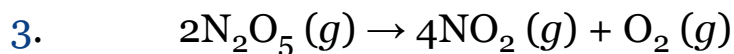
- (d) The overall combustion of butane is an exothermic reaction. Explain why this is, in terms of bond energies.

The overall reaction is exothermic because more energy is released when the products are formed than is required to break the bonds in the reactants.

- (e) One of the major sources of error in this experiment comes from the heat which is absorbed by the air. Why, then, might it not be a good idea to perform this experiment inside a sealed container to prevent the heat from leaving the system?

Performing this reaction in a sealed container would limit the amount of oxygen available to react. To get an accurate heat of combustion for butane, it needs to be the limiting reactant, and if the oxygen runs out prior to all the butane combusting, that will not happen.

Alternatively, the pressure of the sealed container could exceed the strength of the container and cause an explosion. There are approximately twice as many gas molecules after the reaction as there are prior to the reaction, and that means the pressure would approximately double if the reaction goes to completion.



The data below was gathered for the decomposition of N_2O_5 at 310 K via the equation above.

Time (s)	$[\text{N}_2\text{O}_5] (M)$
0	0.250

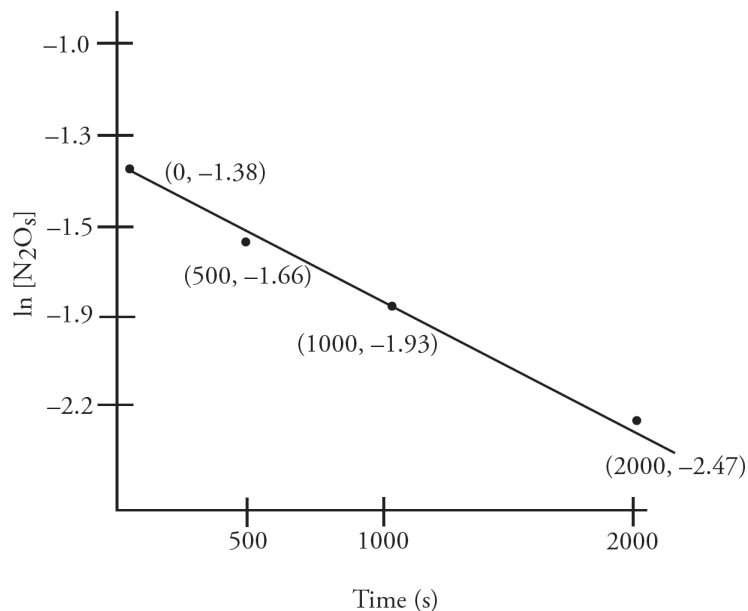
500.	0.190
1000.	0.145
2000.	0.085

- (a) How does the rate of appearance of NO_2 compare to the rate of disappearance of N_2O_5 ? Justify your answer.

Due to the stoichiometric ratios, 4 moles of NO_2 are created for every 2 moles of N_2O_5 that decompose. Therefore, the rate of appearance of NO_2 will be twice the rate of disappearance for N_2O_5 . As that rate is constantly changing over the course of the reaction, it is impossible to get exact values, but the ratio of 2:1 will stay constant.

- (b) The reaction is determined to be first order overall. On the axes below, create a graph of some function of concentration vs. time that will produce a straight line. Label and scale your axes appropriately.

First order decompositions always create a straight line when plotting the natural log of concentration of the reactant vs. time.



(c) (i) What is the rate constant for this reaction? Include units.

Interpreting the graph using slope-intercept form, you get $\ln[\text{N}_2\text{O}_5]_t = -kt + \ln[\text{N}_2\text{O}_5]_0$.

Any (non-intercept) values on the graph can be plugged in to determine the rate constant.

Using at $[\text{N}_2\text{O}_5] = 0.190 \text{ M}$ at $t = 500 \text{ s}$:

$$\ln(0.190) = -k(500.) + \ln(0.250)$$

$$-1.66 = -k(500.) + -1.39$$

$$-.27 = -k(500.)$$

$$k = 5.40 \times 10^{-4}$$

In terms of units, if rate = $k[\text{N}_2\text{O}_5]$, then via analyzing the units:

$$\text{M/s} = k(\text{M}) \quad k = \text{s}^{-1}$$

So $k = 5.40 \times 10^{-4} \text{ s}^{-1}$. Any point along the line will give the same value (as it is equal to the negative slope of the line).

(ii) What would the concentration of N_2O_5 be at $t = 1500$ s?

Using the equation from (i):

$$\ln[\text{N}_2\text{O}_5]_{1500} = -5.40 \times 10^{-4} \text{ s}^{-1}(1500. \text{ s}) + \ln(0.250)$$

$$\ln[\text{N}_2\text{O}_5]_{1500} = -2.20$$

$$[\text{N}_2\text{O}_5]_{1500} = 0.111 \text{ M}$$

(iii) What is the half-life of N_2O_5 ?

The half life is defined as how long it takes half of the original sample to decay. Thus, at the half-life of N_2O_5 for this reaction, $[\text{N}_2\text{O}_5] = 0.125 \text{ M}$. So:

$$\ln(0.125) = -5.40 \times 10^{-4} (t) + \ln(0.250)$$

$$-2.08 = -5.40 \times 10^{-4} (t) + -1.39$$

$$-0.69 = -5.40 \times 10^{-4} (t)$$

$$t = 1280 \text{ s}$$

(d) Would the addition of a catalyst increase, decrease, or have no effect on the following variables? Justify your answers.

(i) Rate of disappearance of N_2O_5

Catalysts speed up the reaction, so adding a catalyst would increase the rate of disappearance of N_2O_5 .

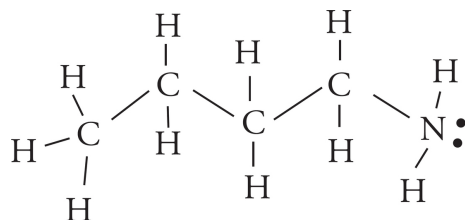
(ii) Magnitude of the rate constant

If the reaction is moving faster, the slope of any line graphing concentration vs. time would be steeper. This increases the magnitude of the slope, and thus, increases the magnitude of the rate constant as well.

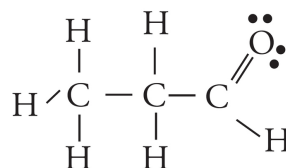
(iii) Half-life of N_2O_5

If the reaction is proceeding at a faster rate, it will take less time for the N_2O_5 to decompose, and thus the half-life of the N_2O_5 would decrease.

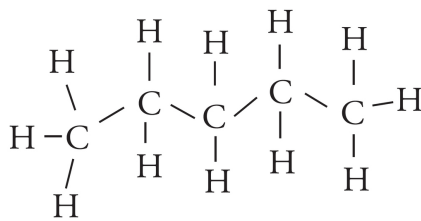
4. Consider the Lewis structures for the following four molecules:



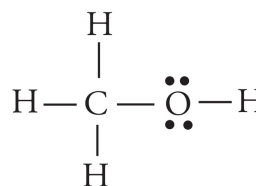
n-Butylamine



Propanal



Pentane



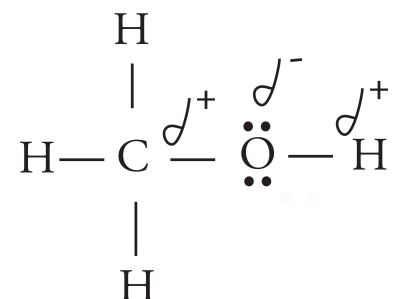
Methanol

- (a) All of the substances are liquids at room temperature. Organize them from high to low in terms of boiling points, clearly differentiating between the intermolecular forces in each substance.

The strongest type of intermolecular force present here is hydrogen bonding, which both n-Butylamine and methanol exhibit. Of the two, n-Butylamine would have the stronger London dispersion forces because it has more electrons (is more polarizable), and so it has stronger IMFs than methanol. For the remaining two structures, propanal has permanent dipoles, while pentane is completely nonpolar. Therefore, propanal would have stronger IMFs than pentane.

So, from high to low boiling point: n-Butylamine > methanol > propanal > pentane.

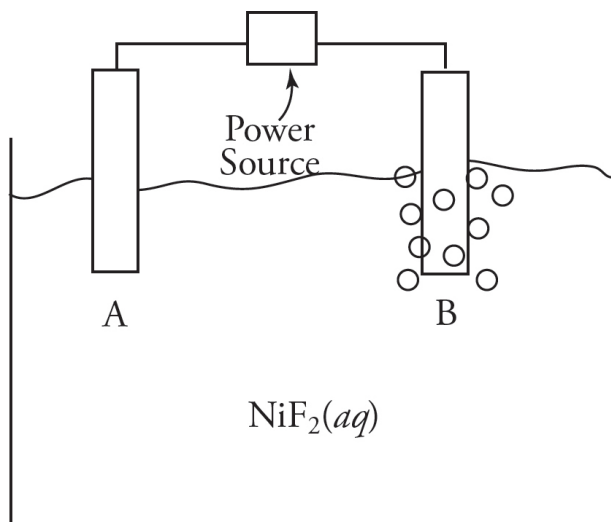
- (b) On the methanol diagram reproduced below, draw the locations of all dipoles.



- (c) n-Butylamine is found to have the lowest vapor pressure at room temperature out of the four liquids. Justify this observation in terms of intermolecular forces.

Vapor pressure arises from molecules overcoming the intermolecular forces to other molecules and escaping the surface of the liquid to become a gas. Due to its hydrogen bonding and large, highly polarizable electron cloud, n-Butylamine would have the strongest intermolecular forces and thus its molecules would have the hardest time escaping the surface, leading to a low vapor pressure.

5. Current is run through an aqueous solution of nickel (II) fluoride, and a gas is evolved at the right-hand electrode, as indicated by the diagram below:



The standard reduction potential for several reactions is given in the following table:

Half-cell	E°_{red}
$\text{F}_2 (g) + 2e^- \rightarrow 2\text{F}^-$	+2.87 V
$\text{O}_2 (g) + 4\text{H}^+ + 4e^- \rightarrow 2\text{H}_2\text{O} (l)$	+1.23 V
$\text{Ni}^{2+} + 2e^- \rightarrow \text{Ni} (s)$	-0.25 V
$2\text{H}_2\text{O} (l) + 2e^- \rightarrow \text{H}_2 (g) + 2\text{OH}^-$	-0.83 V

(a) Determine which half-reaction is occurring at each electrode:

(i) Oxidation:

The two choices for oxidation reductions are $2\text{F}^- \rightarrow \text{F}_2 (g) + 2e^-$ (because fluoride ions are present in the

solution) and $2\text{H}_2\text{O} (l) \rightarrow \text{O}_2 (g) + 4\text{H}^+ + 4e^-$ (as water will lose electrons when oxidized). As you flipped the reactions, you must also flip the signs, so for the fluoride reaction $E^\circ_{\text{ox}} = -2.87 \text{ V}$ and for the water one $E^\circ_{\text{ox}} = -1.23 \text{ V}$. The half-reaction with the less negative value is more likely to occur, so the answer is $2\text{H}_2\text{O} (l) \rightarrow \text{O}_2 (g) + 4\text{H}^+ + 4e^-$.

(ii) Reduction:

Using the same logic as above, the two choices for reduction reaction are $\text{Ni}^{2+} + 2e^- \rightarrow \text{Ni} (s)$ and $2\text{H}_2\text{O} (l) + 2e^- \rightarrow \text{H}_2 (g) + 2\text{OH}^-$. Unlike the oxidation reaction, there is no need to flip the signs on these half-reactions. The more positive value in this case belongs to the nickel reaction, so the answer is $\text{Ni}^{2+} + 2e^- \rightarrow \text{Ni} (s)$.

(b) (i) Calculate the standard cell potential of the cell.

$$E^\circ_{\text{cell}} = E^\circ_{\text{ox}} + E^\circ_{\text{red}} = -1.23 \text{ V} + (-0.25 \text{ V}) = -1.48 \text{ V}$$

(ii) Calculate the Gibbs free energy value for the cell at standard conditions.

$$\Delta G = -nFE$$

n is equal to 4 moles of electrons (from the oxidation reaction). Even though $n = 2$ in the unbalanced reduction reaction, to balance the reaction, the reduction half-reaction would need to be multiplied by 2 so that the electrons balance. Additionally, a volt is equal to a Joule/Coulomb, which is what you should use to get the units to make sense. So:

$$\Delta G = -(4 \text{ mol } e^-)(96,500 \text{ C/mol } e^-)(-0.98 \text{ J/C})$$

$$\Delta G = 380,000 \text{ J or } 380 \text{ kJ}$$

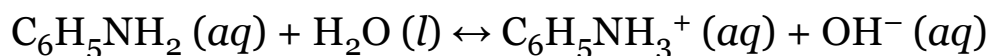
- (c) Which electrode in the diagram (A or B) is the cathode, and which is the anode? Justify your answers.

Oxygen gas is evolved in the oxidation reaction, meaning that is occurring at electrode B. Oxidation always occurs at the anode, so electrode A is the cathode and electrode B is the anode.

6. Aniline, $\text{C}_6\text{H}_5\text{NH}_2$, is a weak base with $K_b = 3.8 \times 10^{-10}$.

- (a) Write out the reaction that occurs when aniline reacts with water.

The aniline will act as a proton acceptor and take a proton from the water.



- (b) (i) What is the concentration of each species at equilibrium in a solution of $0.25 \text{ M } \text{C}_6\text{H}_5\text{NH}_2$?

For the above equilibrium, $K_b = [\text{C}_6\text{H}_5\text{NH}_3^+][\text{OH}^-]/[\text{C}_6\text{H}_5\text{NH}_2]$. The concentrations of both the conjugate acid and the hydroxide ion will be equal, and the concentration of the aniline itself will be approximately the same, as it is a weak base which has a very low protonation rate. You can do an ICE chart to confirm this, but it is not really necessary if you understand the concepts underlying weak acids and bases.

$$3.8 \times 10^{-10} = (x)(x)/(0.25)$$

$$x^2 = 9.5 \times 10^{-11}$$

$$x = 9.7 \times 10^{-6}$$

$$[\text{C}_6\text{H}_5\text{NH}_3^+] = [\text{OH}^-] = 9.7 \times 10^{-6} \text{ M} \text{ and } [\text{C}_6\text{H}_5\text{NH}_2] = 0.25 \text{ M}$$

(ii) What is the pH value for the solution in (b)(i)?

$$\text{pOH} = -\log[\text{OH}^-]$$

$$\text{pOH} = -\log(9.7 \times 10^{-6} \text{ M})$$

$$\text{pOH} = 5.01$$

$$\text{pOH} + \text{pH} = 14$$

$$5.01 + \text{pH} = 14$$

$$\text{pH} = 8.99$$

7. A rigid, sealed 12.00 L container is filled with 10.00 g each of three different gases: CO_2 , NO, and NH_3 . The temperature of the gases is held constant 35.0°C . Assume ideal behavior for all gases.

(a) (i) What is the mole fraction of each gas?

First, the moles of each gas need calculating:

$$\text{CO}_2 = 10.00 \text{ g} \times 1 \text{ mol}/44.01 \text{ g} = 0.227 \text{ mol}$$

$$\text{NO} = 10.00 \text{ g} \times 1 \text{ mol}/30.01 \text{ g} = 0.333 \text{ mol}$$

$$\text{NH}_3 = 10.00 \text{ g} \times 1 \text{ mol}/17.03 \text{ g} = 0.587 \text{ mol}$$

$$\text{Total moles of gas} = 1.147 \text{ moles}$$

$$X_{\text{CO}_2} = 0.227/1.147 = 0.198$$

$$X_{\text{NO}} = 0.333/1.147 = 0.290$$

$$X_{\text{NH}_3} = 0.587/1.147 = 0.512$$

(ii) What is the partial pressure of each gas?

Using the Ideal Gas Law, we can calculate the total pressure in the container:

$$PV = nRT$$

$$P(12.00\text{ L}) = (1.147\text{ mol})(0.0821\text{ atm}\times\text{L/mol}\cdot\text{K})(308\text{ K})$$

$$P = 2.42\text{ atm}$$

The partial pressure of a gas is equal to the total pressure times the mole fraction of that gas.

$$P_{\text{CO}_2} = (2.42\text{ atm})(0.198) = 0.479\text{ atm}$$

$$P_{\text{NO}} = (2.42\text{ atm})(0.290) = 0.702\text{ atm}$$

$$P_{\text{NH}_3} = (2.42\text{ atm})(0.512) = 1.24\text{ atm}$$

- (b) Out of the three gases, molecules of which gas will have the highest velocity? Why?

If all gases are at the same temperature, they have the same amount of kinetic energy. KE has aspects of both mass and velocity, so the gas with the lowest mass would have the highest velocity. Thus, the NH_3 molecules have the highest velocity.

- (c) Name one circumstance in which the gases might deviate from ideal behavior, and clearly explain the reason for the deviation.

The most common reason for deviation from ideal behavior is that the intermolecular forces of the gas molecules are acting upon one another. This would occur when the molecules are very close together and/or moving very slowly, so deviations would occur at high pressures and/or low temperatures.

Part VII

Additional Practice Tests

Practice Test 3

Click [here](#) to download and print the PDF version of this exercise.

AP[®] Chemistry Exam

SECTION I: Multiple-Choice Questions

DO NOT OPEN THIS BOOKLET UNTIL YOU ARE TOLD TO DO SO.

At a Glance

Total Time

1 hour and 30 minutes

Number of Questions

60

Percent of Total Grade

50%

Writing Instrument

Pencil required

Instructions

Section I of this examination contains 60 multiple-choice questions. Fill in only the ovals for numbers 1 through 60 on your answer sheet.

CALCULATORS MAY NOT BE USED IN THIS PART OF THE EXAMINATION.

Indicate all of your answers to the multiple-choice questions on the answer sheet. No credit will be given for anything written in this exam booklet, but you may use the booklet for notes or scratch work.

After you have decided which of the suggested answers is best, completely fill in the corresponding oval on the answer sheet. Give only one answer to each question. If you change an answer, be sure that the previous mark is erased completely. Here is a sample question and answer.

Sample Question

Chicago is a

- (A) state
- (B) city
- (C) country
- (D) continent

Sample Answer

☐ (A) ☒ (B) ☐ (C) ☐ (D)

Use your time effectively, working as quickly as you can without losing accuracy. Do not spend too much time on any one question. Go on to other questions and come back to the ones you have not answered if you have time. It is not expected that everyone will know the answers to all the multiple-choice questions.

About Guessing

Many candidates wonder whether or not to guess the answers to questions about which they are not certain. Multiple-choice scores are based on the number of questions answered correctly. Points are not deducted for incorrect answers, and no points are awarded for unanswered questions. Because points are not deducted for incorrect answers, you are encouraged to answer all multiple-choice questions. On any questions you do not know the answer to, you should eliminate as many choices as you can, and then select the best answer among the remaining choices.

CHEMISTRY

SECTION I

Time—1 hour and 30 minutes

INFORMATION IN THE TABLE BELOW AND ON THE FOLLOWING PAGES MAY BE USEFUL IN ANSWERING THE QUESTIONS IN THIS SECTION OF THE EXAMINATION.

DO NOT DETACH FROM BOOK.

PERIODIC TABLE OF THE ELEMENTS

1 H 1.008																	2 He 4.00
3 Li 6.94	4 Be 9.01											5 B 10.81	6 C 12.01	7 N 14.01	8 O 16.00	9 F 19.00	10 Ne 20.18
11 Na 22.99	12 Mg 24.30											13 Al 26.98	14 Si 28.09	15 P 30.97	16 S 32.06	17 Cl 35.45	18 Ar 39.95
19 K 39.10	20 Ca 40.08	21 Sc 44.96	22 Ti 47.90	23 V 50.94	24 Cr 52.00	25 Mn 54.94	26 Fe 55.85	27 Co 58.93	28 Ni 58.69	29 Cu 63.55	30 Zn 65.39	31 Ga 69.72	32 Ge 72.59	33 As 74.92	34 Se 78.96	35 Br 79.90	36 Kr 83.80
37 Rb 85.47	38 Sr 87.62	39 Y 88.91	40 Zr 91.22	41 Nb 92.91	42 Mo 95.94	43 Tc (98)	44 Ru 101.1	45 Rh 102.91	46 Pd 106.42	47 Ag 107.87	48 Cd 112.41	49 In 114.82	50 Sn 118.71	51 Sb 121.75	52 Te 127.60	53 I 126.91	54 Xe 131.29
55 Cs 132.91	56 Ba 137.33	57 *La 138.91	72 Hf 178.49	73 Ta 180.95	74 W 183.85	75 Re 186.21	76 Os 190.2	77 Ir 192.2	78 Pt 195.08	79 Au 196.97	80 Hg 200.59	81 Tl 204.38	82 Pb 207.2	83 Bi 208.98	84 Po (209)	85 At (210)	86 Rn (222)
87 Fr (223)	88 Ra 226.02	89 †Ac 227.03	104 Rf (261)	105 Db (262)	106 Sg (266)	107 Bh (264)	108 Hs (277)	109 Mt (268)	110 Ds (271)	111 Rg (272)							

*Lanthanide Series

58 Ce 140.12	59 Pr 140.91	60 Nd 144.24	61 Pm (145)	62 Sm 150.4	63 Eu 151.97	64 Gd 157.25	65 Tb 158.93	66 Dy 162.50	67 Ho 164.93	68 Er 167.26	69 Tm 168.93	70 Yb 173.04	71 Lu 174.97
90 Th 232.04	91 Pa 231.04	92 U 238.03	93 Np (237)	94 Pu (244)	95 Am (243)	96 Cm (247)	97 Bk (247)	98 Cf (251)	99 Es (252)	100 Fm (257)	101 Md (258)	102 No (259)	103 Lr (262)

†Actinide Series

ADVANCED PLACEMENT CHEMISTRY EQUATIONS AND CONSTANTS

Throughout the test the following symbols have the definitions specified unless otherwise noted.

L, mL = liter(s), milliliter(s)
 g = gram(s)
 nm = nanometer(s)
 atm = atmosphere(s)

mm Hg = millimeters of mercury
 J, kJ = joule(s), kilojoule(s)
 V = volt(s)
 mol = mole(s)

ATOMIC STRUCTURE

$$E = h\nu$$

$$c = \lambda\nu$$

E = energy
 ν = frequency
 λ = wavelength

Planck's constant, $h = 6.626 \times 10^{-34}$ J s

Speed of light, $c = 2.998 \times 10^8$ m s⁻¹

Avogadro's number = 6.022×10^{23} mol⁻¹

Electron charge, $e = -1.602 \times 10^{-19}$ coulomb

EQUILIBRIUM

$$K_c = \frac{[C]^c[D]^d}{[A]^a[B]^b}, \text{ where } aA + bB \rightleftharpoons cC + dD$$

$$K_p = \frac{(P_C)^c(P_D)^d}{(P_A)^a(P_B)^b}$$

$$K_a = \frac{[H^+][A^-]}{[HA]}$$

$$K_b = \frac{[OH^-][HB^+]}{[B]}$$

$$K_w = [H^+][OH^-] = 1.0 \times 10^{-14} \text{ at } 25^\circ\text{C}$$

$$= K_a \times K_b$$

$$\text{pH} = -\log[H^+], \text{ pOH} = -\log[OH^-]$$

$$14 = \text{pH} + \text{pOH}$$

$$\text{pH} = \text{p}K_a + \log \frac{[A^-]}{[HA]}$$

$$\text{p}K_a = -\log K_a, \text{ p}K_b = -\log K_b$$

Equilibrium Constants

K_c (molar concentrations)

K_p (gas pressures)

K_a (weak acid)

K_b (weak base)

K_w (water)

KINETICS

$$\ln[A]_t - \ln[A]_0 = -kt$$

$$\frac{1}{[A]_t} - \frac{1}{[A]_0} = kt$$

$$t_{1/2} = \frac{0.693}{k}$$

k = rate constant

t = time

$t_{1/2}$ = half-life

GASES, LIQUIDS, AND SOLUTIONS

$$PV = nRT$$

$$P_A = P_{\text{total}} \times X_A, \text{ where } X_A = \frac{\text{moles A}}{\text{total moles}}$$

$$P_{\text{total}} = P_A + P_B + P_C + \dots$$

$$n = \frac{m}{M}$$

$$K = ^\circ\text{C} + 273$$

$$D = \frac{m}{V}$$

$$KE \text{ per molecule} = \frac{1}{2}mv^2$$

Molarity, M = moles of solute per liter of solution

$$A = abc$$

P = pressure

V = volume

T = temperature

n = number of moles

m = mass

M = molar mass

D = density

KE = kinetic energy

v = velocity

A = absorbance

a = molar absorptivity

b = path length

c = concentration

Gas constant, $R = 8.314 \text{ J mol}^{-1} \text{ K}^{-1}$

$$= 0.08206 \text{ L atm mol}^{-1} \text{ K}^{-1}$$

$$= 62.36 \text{ L torr mol}^{-1} \text{ K}^{-1}$$

$$1 \text{ atm} = 760 \text{ mm Hg}$$

$$= 760 \text{ torr}$$

STP = 273.15 K and 1.0 atm

Ideal gas at STP = 22.4 L mol⁻¹

THERMOCHEMISTRY/ ELECTROCHEMISTRY

$$q = mc\Delta T$$

$$\Delta S^\circ = \sum S^\circ \text{ products} - \sum S^\circ \text{ reactants}$$

$$\Delta H^\circ = \sum \Delta H_f^\circ \text{ products} - \sum \Delta H_f^\circ \text{ reactants}$$

$$\Delta G^\circ = \sum \Delta G_f^\circ \text{ products} - \sum \Delta G_f^\circ \text{ reactants}$$

$$\Delta G^\circ = \Delta H^\circ - T\Delta S^\circ$$

$$= -RT \ln K$$

$$= -nFE^\circ$$

$$I = \frac{q}{t}$$

q = heat

m = mass

c = specific heat capacity

T = temperature

S° = standard entropy

H° = standard enthalpy

G° = standard free energy

n = number of moles

E° = standard reduction potential

I = current (amperes)

q = charge (coulombs)

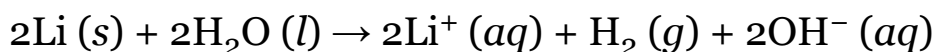
t = time (seconds)

Faraday's constant, $F = 96,485$ coulombs per mole of electrons

$$1 \text{ volt} = \frac{1 \text{ joule}}{1 \text{ coulomb}}$$

Acid	Lewis Diagram	K_a
HOBr	$\begin{array}{c} \cdot\cdot \quad \cdot\cdot \\ \text{H}-\text{O}-\text{Br}: \\ \cdot\cdot \quad \cdot\cdot \end{array}$	2.0×10^{-9}
HOCl	$\begin{array}{c} \cdot\cdot \quad \cdot\cdot \\ \text{H}-\text{O}-\text{Cl}: \\ \cdot\cdot \quad \cdot\cdot \end{array}$	3.0×10^{-8}

- The K_a values and Lewis structures for two different oxoacids are listed above. Which of the following statements correctly identifies the stronger acid and offers up the best justification?
 - HOBr, because bromine is larger than chlorine and strongly repels the $-\text{OH}$ group
 - HOBr, because bromine's electron cloud is more polarizable than chlorine's
 - HOCl, because the OCl^- ion has a stronger attraction for protons than the OBr^- ion
 - HOCl, because chlorine is more electronegative than bromine and weakens the $\text{H}-\text{O}$ bond more effectively
- Approximately how many grams of oxygen are present in a 10.0 g sample of NaOH?
 - 2.00 g
 - 4.00 g
 - 6.00 g
 - 8.00 g



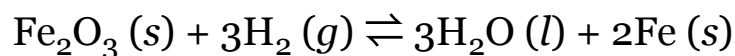
- A 3.50 g sample of lithium metal is dropped into a beaker of water, causing the above reaction to occur. How many liters of hydrogen gas are produced if this reaction takes place at STP?

- (A) 5.60 L
- (B) 11.2 L
- (C) 22.4 L
- (D) 28.0 L

Metal Initial Temperature	98.0°C
Water Initial Temperature	26.0°C
Water + Metal Final Temperature	28.0°C
Mass Metal	10.0 g
Mass Water	100. g

4. A piece of an unknown metal is heated to a high temperature, and then dropped into a cup of water. Given the data above and assuming no heat is lost to the environment, what is the approximate specific heat of the metal? (Water has a specific heat of 4.2 J/g°C.)

- (A) 0.60 J/g°C
- (B) 0.80 J/g°C
- (C) 1.00 J/g°C
- (D) 1.20 J/g°C



5. What would the equilibrium constant expression, K_c , be for the above reaction?

(A) $K_c = \frac{[\text{Fe}]^2 [\text{H}_2\text{O}]^3}{[\text{Fe}_2\text{O}_3] [\text{H}_2]^3}$

$$(B) K_c = \frac{[H_2O]^3}{[H_2]^3}$$

$$(C) K_c = \frac{[H_2]^3}{[H_2O]^3}$$

$$(D) K_c = \frac{1}{[H_2]^3}$$

Questions 6-10 refer to the following standard reduction potentials.

Half Reaction	Reduction Potential
$O_2 (g) + 4H^+ (aq) + 4e^- \rightarrow 2H_2O (l)$	1.23 V
$Ag^+ (aq) + e^- \rightarrow Ag (s)$	0.80 V
$Cu^{2+} (aq) + 2e^- \rightarrow Cu (s)$	0.34 V
$Fe^{2+} (aq) + 2e^- \rightarrow Fe (s)$	-0.44 V
$2H_2O (l) + 2e^- \rightarrow H_2 (g) + 2OH^-$	-0.83 V
$Al^{3+} (aq) + 3e^- \rightarrow Al (s)$	-1.66 V

6. What would the cell potential be for a galvanic cell constructed of Ag/Ag^+ and Fe/Fe^{2+} half-cells?

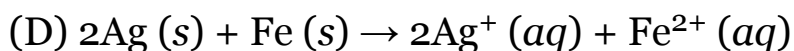
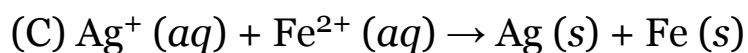
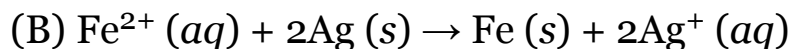
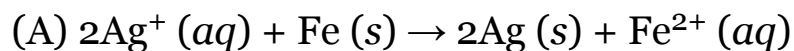
(A) 0.36 V

(B) 1.16 V

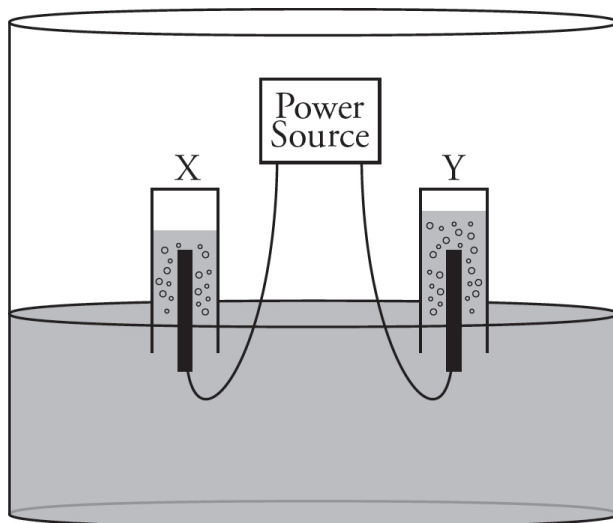
(C) 1.24 V

(D) 2.04 V

7. What would the net ionic reaction be for the galvanic cell described in question #6?



8. A 2.0 A current is applied to beakers containing identical volumes of each of the following 1.0 M solutions. In beaker will the greatest mass of metal plate out?



9. A current is run through a solution of pure water, and gases are collected as shown in the above diagram. Which of the following correctly identifies the cathode and anode during the hydrolysis, along with identifying the gas present at each?

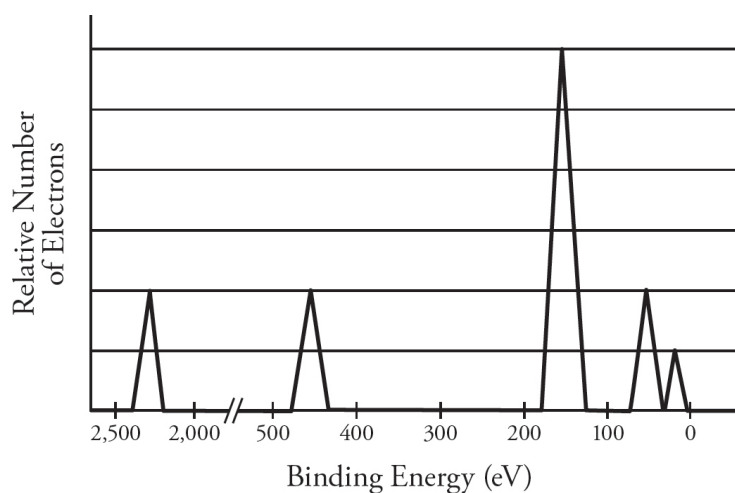
	Electrode X Identity	Gas collected at Electrode X

(A)	Cathode	Hydrogen
(B)	Anode	Hydrogen
(C)	Cathode	Oxygen
(D)	Anode	Oxygen

10. Which of the following would NOT cause a reaction to occur?

- (A) Adding some aluminum pellets to a solution containing Fe^{2+} ions
- (B) Placing a copper strip into a solution containing Ag^+ ions
- (C) Adding powdered iron to a solution containing Cu^{2+} ions
- (D) Dropping a silver coin into a solution containing Al^{3+} ions

Use the following information to answer questions 11-13.



Peak 1	Peak 2	Peak 3	Peak 4	Peak 5
2300 eV	450 eV	150 eV	30 eV	5.0 eV

The photoelectron spectrum for a neutral aluminum atom is located above.

11. The amount of energy necessary to remove an electron from the $2p$ subshell is closest to which value?

- (A) 450 eV
- (B) 150 eV
- (C) 30 eV
- (D) 5.0 eV

12. On a spectrum of an aluminum ion:

- (A) All peaks would be identical.
- (B) The peak furthest to the right be twice as tall.
- (C) The two peaks furthest to the right would be missing.
- (D) All peaks would be half as tall.

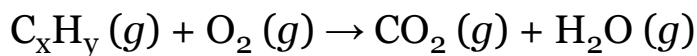
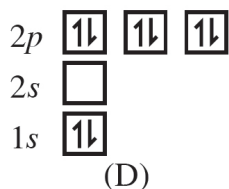
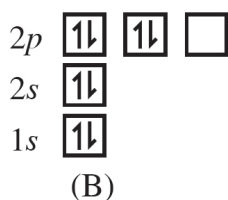
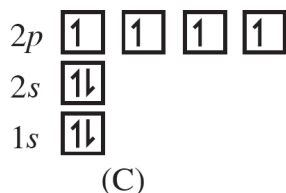
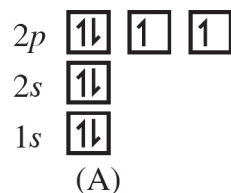
13. A different aluminum atom is exposed to incoming radiation with an energy of 200 eV. Ejected electrons that were originally in which orbital would have the lowest kinetic energy?

- (A) $1s$
- (B) $2s$
- (C) $2p$
- (D) $3p$

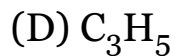
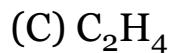
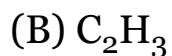
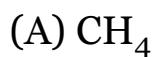
14. A current of 4.0 A is run through an electrolytic cell for a total of 2.0 minutes. How many coulombs of charge passed through the cell during its operation?

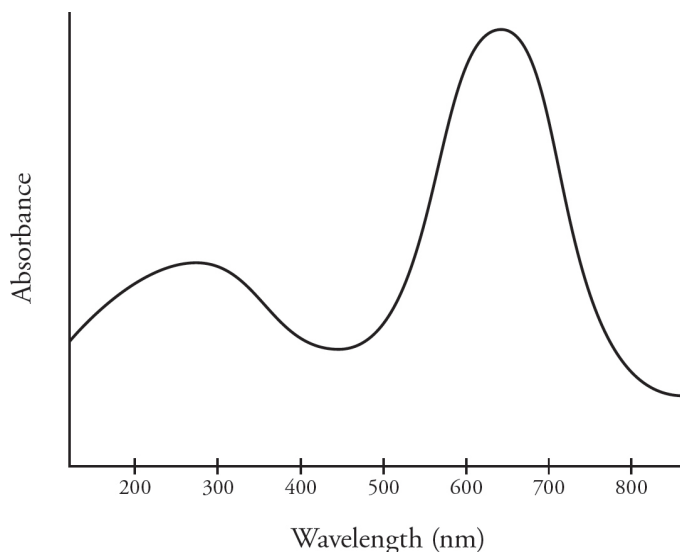
- (A) 2.0 C
- (B) 8.0 C
- (C) 30 C
- (D) 480 C

15. Which of the following orbital diagrams shows the correct electron configuration for an oxygen atom?



16. A hydrocarbon of an unknown formula consisting of only hydrogen and carbon is combusted in excess oxygen, producing 44.0 g of carbon dioxide and 18.0 g of water via the above **unbalanced** reaction. Which of the following is a potential formula for the hydrocarbon?

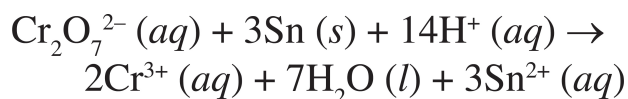




17. Cu^{2+} ions appear blue in solution, and the NO_3^- ion is colorless. The absorbance of a $\text{Cu}(\text{NO}_3)_2$ solution is measured at various wavelengths using a spectrophotometer. To determine the concentration of a $\text{Cu}(\text{NO}_3)_2$ solution of unknown concentration, what wavelength of light would produce the best results?

- (A) 350 nm
- (B) 500 nm
- (C) 640 nm
- (D) 800 nm

Use the following information to answer questions 18-20.



A solution of potassium dichromate, $\text{K}_2\text{Cr}_2\text{O}_7$, is poured over a piece of solid tin, causing the oxidation-reduction reaction above to occur.

18. Which substance is oxidized, and which is reduced?

	Oxidized	Reduced
(A)	Cr_2O_7	Sn
(B)	H^+	Cr_2O_7
(C)	Sn	Cr_2O_7
(D)	Cr_2O_7	H^+

19. How many moles of electrons are transferred in the balanced equation?

- (A) 2
- (B) 3
- (C) 4
- (D) 6

20. How many grams of tin will react with 100. mL of 0.033 M $\text{K}_2\text{Cr}_2\text{O}_7$?

- (A) 0.40 g
- (B) 1.2 g
- (C) 2.4 g
- (D) 3.6 g

Salt	ΔH_{soln} (kJ/mol)
LiClO_4	-26.55
NaClO_4	13.88
KClO_4	41.38

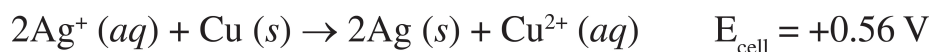
21. The enthalpy of solution for three different perchlorate compounds is listed in the table above. Based on the data, which of the following statements is correct?

- (A) As cation size increases, the rate at which the lattice energy is changing exceeds the rate at which the hydration energy is changing.
- (B) The smaller the cation, the smaller the magnitude of the lattice energy will be.
- (C) As cation size increases, the magnitude of the temperature change that occurs in the surrounding solution consistently increases.
- (D) Of the three salts, only lithium perchlorate is fully soluble in water, which is demonstrated by its negative enthalpy of solution.



22. The above system is at equilibrium. Which of the following would cause an immediate increase in the rate of the reverse reaction?

- (A) Adding some water
- (B) Adding some $\text{AuNO}_3 (aq)$
- (C) Increasing the pressure
- (D) Increasing the temperature



23. A silver/copper galvanic cell is set up, and the above reaction occurs. If all other variables were to remain the same, what is one way to increase the cell potential?

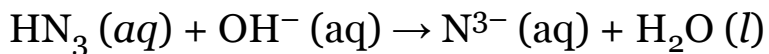
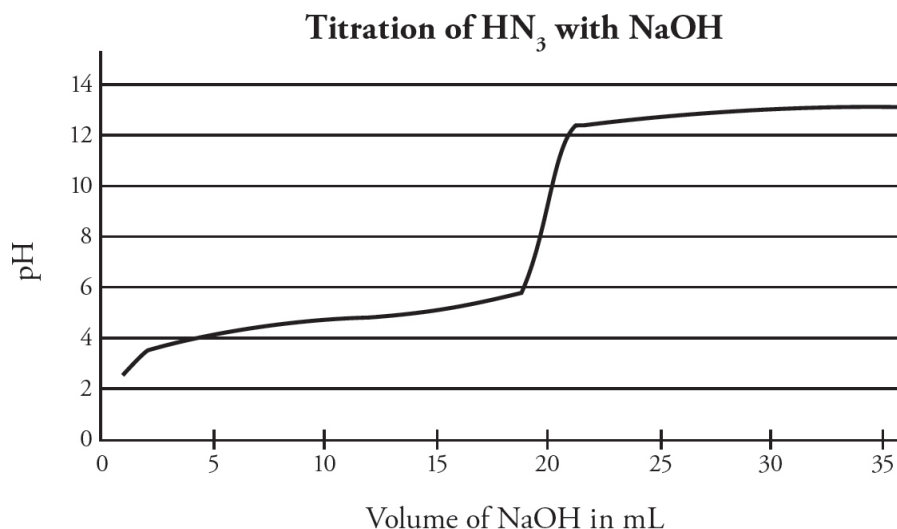
- (A) Increase the mass of the copper electrode

(B) Increase the mass of the silver electrode

(C) Increase $[\text{Ag}^+]$

(D) Increase $[\text{Cu}^{2+}]$

Use the following information to answer questions 24-27.



25.0 mL of hydrazoic acid, HN_3 ($K_a = 1.9 \times 10^{-5}$), is titrated with 0.50 M NaOH . The pH is measured during the titration, producing the above graph. The equivalence point is reached at a volume of 20.0 mL of NaOH added.

24. What is the concentration of the HN_3 ?

(A) 0.40 M

(B) 0.50 M

(C) 0.60 M

(D) 0.80 M

25. At what volume of NaOH added would $[\text{N}_3^-] = [\text{HN}_3]$?

- (A) 5.0 mL
- (B) 10.0 mL
- (C) 20.0 mL
- (D) 25.0 mL

26. Which indicator would be the best to use to determine the endpoint of the titration?

	Indicator	p<i>K</i>_a
(A)	Methyl Violet	0.80
(B)	Congo Red	4.0
(C)	Thymol Blue	8.9
(D)	Indigo Carmine	12.2

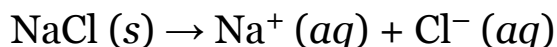
27. If the hydrazoic acid were replaced with hydrocyanic acid ($K_a = 6.2 \times 10^{-10}$) of identical concentration, which of the following changes would occur?

- (A) Less NaOH would be required to reach the titration endpoint.
- (B) The initial pH would change as soon as any NaOH is added will be greater.
- (C) The pH of the solution at the 30.0 mL mark would be higher.
- (D) The pH at the equivalence point would be decreased.

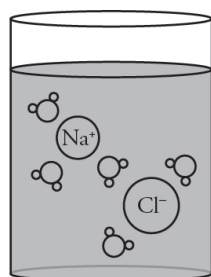
Ion	Ionization energy (kJ/mol)
Al ³⁺	11500
Mg ²⁺	7732
Na ⁺	4562

28. The next ionization energy values for different ions of three period 3 metals are listed above. The reason that the aluminum ion has the greatest next ionization energy is:

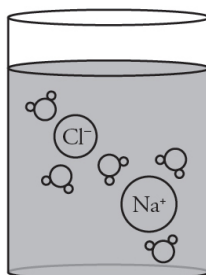
- (A) Aluminum metal is less chemically active than magnesium or sodium.
- (B) The radius of the aluminum ion is the smallest.
- (C) The aluminum ion has the greatest number of unpaired electrons.
- (D) The aluminum ion experiences the smallest amount of shielding.



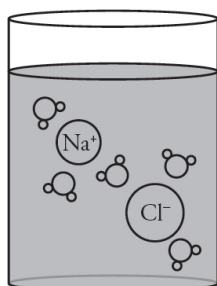
29. A sample of sodium chloride, NaCl, is dissolved fully in water. Which diagram below is an accurate representation of how the solution would look on the particulate level?



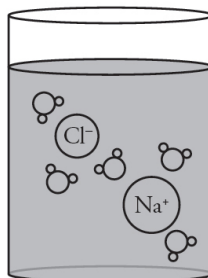
(A)



(B)



(C)



(D)

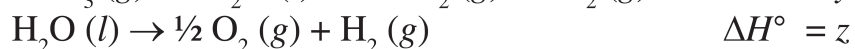
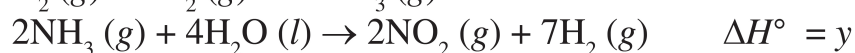
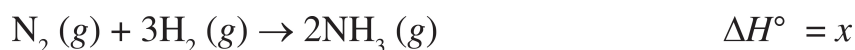
30. Which of the following expressions would accurately give the magnitude for the density of a sample of helium gas at STP?

(A) $\frac{(4.0)}{(0.0821)(273)}$

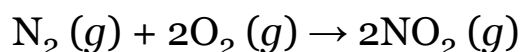
(B) $\frac{(273)(0.0821)}{4.0}$

(C) $\frac{(273)(4.0)}{(0.0821)}$

(D) $(4.0)(273)(0.0821)$



31. Given the enthalpy changes for the above reactions, what would be the enthalpy change for the following reaction?

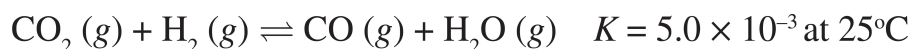


(A) $x + 2y + 2z$

(B) $4x - y - z$

(C) $x + y - 4z$

(D) $2x + y - z$



32. Using the above information, determine the equilibrium constant for $2\text{CO}(g) + 2\text{H}_2\text{O}(g) \rightleftharpoons 2\text{CO}_2(g) + 2\text{H}_2(g)$.

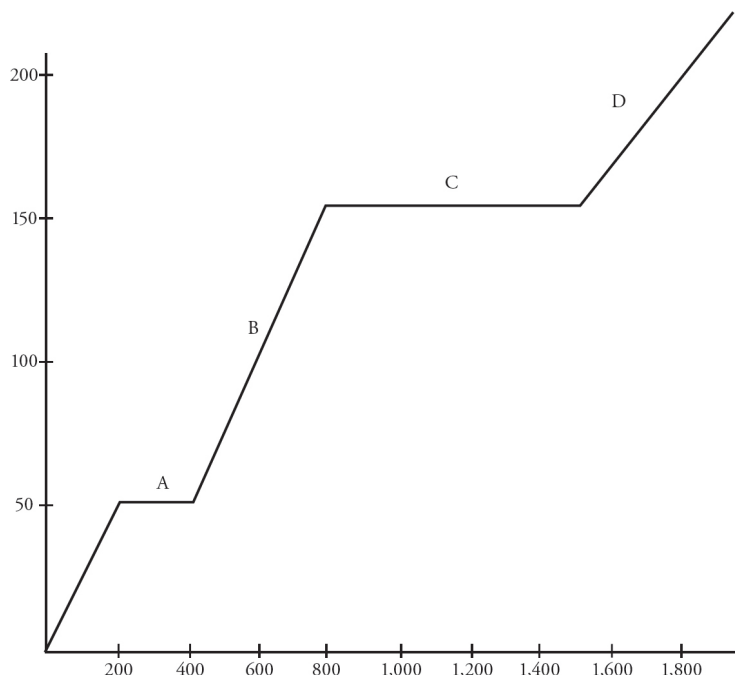
(A) -1.0×10^{-4}

(B) 2.0×10^3

(C) 1.0×10^4

(D) 4.0×10^4

Use the following information to answer questions 33-37.



A 2.0 g sample of a solid covalent substance is placed inside an enclosed chamber, and heat is added at a constant rate. The temperature of the substance is tracked as heat is added, producing the above graph.

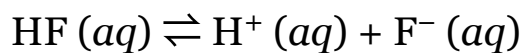
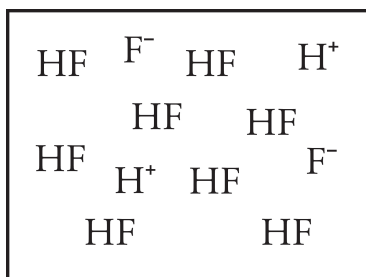
33. At which point on the graph will the substance exist in both its solid and liquid phase simultaneously?

- (A) Point A
- (B) Point B
- (C) Point C
- (D) Point D

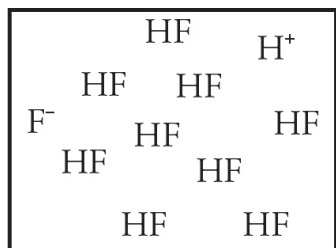
34. At which point(s) on the graph will intermolecular forces be the weakest?”

- (A) Point A only
- (B) Points B and D

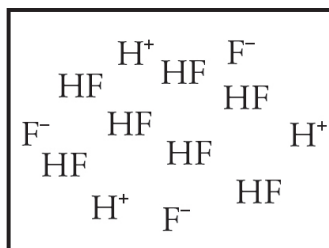
- (C) Point D only
- (D) Intermolecular forces are present at all times.
35. Which of the following statements is true, according to the graph?
- (A) The intramolecular forces are strongest during phase A.
- (B) The heat of vaporization exceeds the heat of fusion.
- (C) The strength of the intermolecular forces is unaffected by the addition of energy.
- (D) The average molecular speed increases throughout the heating process.
36. How much energy would be required to melt a separate 1.0 g sample of the substance?
- (A) 100 J
- (B) 200 J
- (C) 300 J
- (D) 400 J
37. What is the approximate specific heat of the substance in its liquid phase?
- (A) 2.0 J/g^{°C}
- (B) 4.0 J/g^{°C}
- (C) 6.0 J/g^{°C}
- (D) 8.0 J/g^{°C}



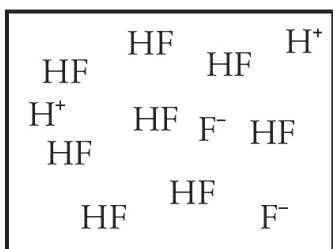
38. HF, is a weak acid with a K_a value of 7.2×10^{-4} . The box above represents a particulate-level view of a HF solution of known concentration. If some water is added, which of the following options would represent a particulate representation of the diluted solution?



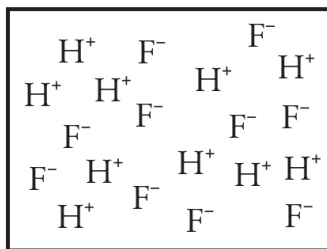
(A)



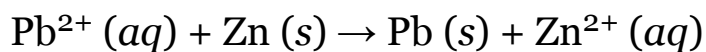
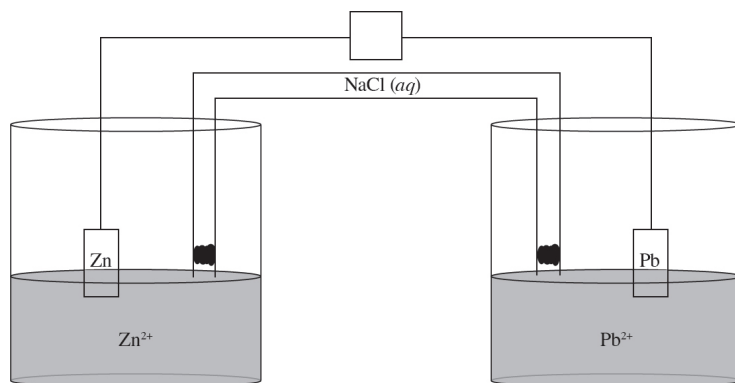
(C)



(B)



(D)



	$[\text{Pb}^{2+}]$	$[\text{Zn}^{2+}]$	Mass Pb electrode	Mass Zn electrode
Battery X	1.0 M	1.0 M	10. g	10. g
Battery Y	2.0 M	2.0 M	20. g	20. g

39. Two batteries are constructed via the above diagram. The concentration of the solutions and the mass of the electrodes are identified in the data table below the diagram. How would the voltage and the battery life for the two batteries compare?

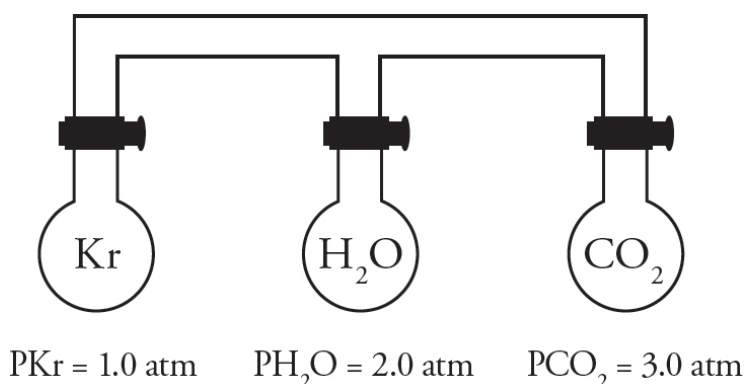
	Voltage	Battery Life
(A)	$X < Y$	$X < Y$
(B)	$X < Y$	$X = Y$
(C)	$X = Y$	$X = Y$
(D)	$X = Y$	$X < Y$

40. Which of the four substances listed below would have the highest melting point?

(A) KCl

- (B) CaBr_2
- (C) MgO
- (D) $\text{C}_6\text{H}_{12}\text{O}_6$

Use the following information to answer questions 41-43.



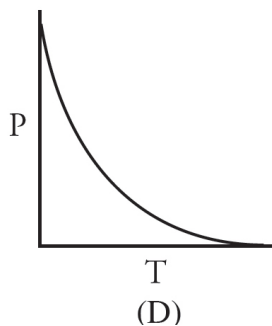
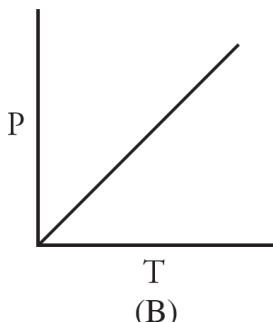
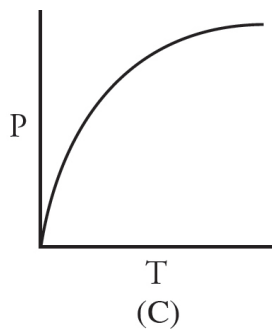
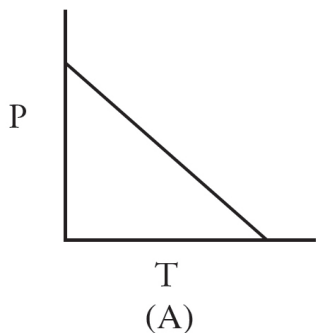
Three identical 1.0 L containers filled with different gases are connected using glass tubes. The stopcocks at the top of each flask are initially closed and all three containers are held at 125°C .

41. Which gas particles are moving the fastest?
- (A) Kr
 - (B) H_2O
 - (C) CO_2
 - (D) All gas particles are moving at the same speed
42. Assuming the volume of the connecting tubes is negligible and that temperature remains constant, what would the total pressure of the system be after all three stopcocks are opened and the gases fully mix?
- (A) 2.0 atm
 - (B) 4.0 atm

(C) 6.0 atm

(D) 8.0 atm

43. After the stopcocks have been opened, the entire system is heated. Choose the graph below that correctly shows how the pressure would change as the temperature is increased.



44. A 1.0 M solution of which of the following would have the highest conductivity?

(A) $\text{C}_2\text{H}_5\text{OH} (aq)$

(B) $\text{NH}_4\text{Cl} (aq)$

(C) $\text{PF}_3 (aq)$

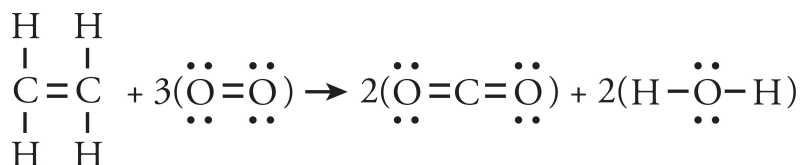
(D) $\text{Na}_2\text{CO}_3 (aq)$

Halogen	Boiling Point ($^{\circ}\text{C}$)
F_2	-188

Cl ₂	-35
Br ₂	59
I ₂	184

45. The boiling point trend observed in the halogens is due to which of the following trends increasing going down the group?

- (A) Molar mass
- (B) First ionization energy
- (C) Electron cloud polarizability
- (D) Electronegativity



Bond	Enthalpy (kJ/mol)
C-H	410
C=C	720
O=O	500
C=O	800
H-O	470

46. Given the bond enthalpy values in the data table above, determine the enthalpy change for the reaction above it.

- (A) -1220 kJ/mol
- (B) - 400 kJ/mol
- (C) 400 kJ/mol

(D) 1220 kJ/mol



47. At 0°C, the partial pressure of each gas above in an equilibrium system are $P_{\text{NO}_2} = 0.60$ atm, $P_{\text{NO}} = 0.30$ atm, and $P_{\text{O}_2} = 0.20$ atm. What is K_p for this reaction at 0°C?

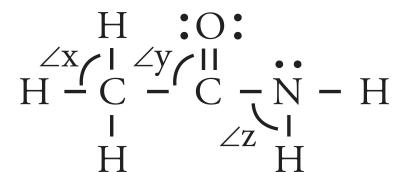
(A) 0.05

(B) 0.10

(C) 10.

(D) 20.

Use the following information to answer questions 48-50.



A Lewis diagram of the acetamide molecule, CH_3CONH_2 , is drawn above. Note the bond angles are not drawn to scale.

48. Identify the bond angles on the diagram from greatest to smallest.

(A) $X > Y > Z$

(B) $X > Z > Y$

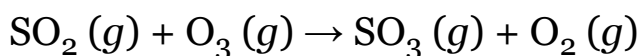
(C) $Z > X > Y$

(D) $Y > X > Z$

49. How many pi bonds are present in an acetamide molecule?

(A) 0

- (B) 1
(C) 2
(D) 3
50. What is the strongest type of intermolecular force present between acetamide molecules in an aqueous solution?
- (A) London dispersion forces
(B) Hydrogen bonding
(C) Ionic bonding
(D) Covalent bonding
51. Chlorous acid, HClO_2 , has a K_a value of 1.0×10^{-2} . What would the pH of a solution of 1.0 M sodium chlorite, NaClO_2 , be?
- (A) 4
(B) 6
(C) 8
(D) 10



Trial	$[\text{SO}_2]$ (M)	$[\text{O}_3]$ (M)	Initial Rate (M/s)
1	0.10	0.10	0.075
2	0.10	0.20	0.075
3	0.20	0.10	0.300

52. The above reaction is run three separate times, varying the initial concentration of both reactants as shown. The initial rate

is shown for each reaction. What is the rate law for the reaction?

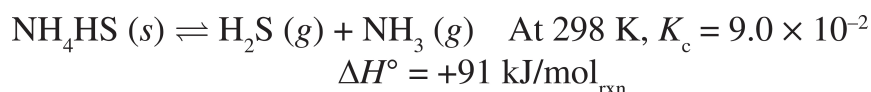
(A) $\text{Rate} = k[\text{SO}_2][\text{O}_3]$

(B) $\text{Rate} = k[\text{SO}_2]$

(C) $\text{Rate} = k[\text{SO}_2]^2$

(D) $\text{Rate} = k[\text{O}_3]^2$

Use the following information to answer questions 53-56.



Some solid NH_4HS is placed in a sealed and evacuated 250 mL flask at 298 K, and the above decomposition reaction is allowed to reach equilibrium.

53. Calculate the concentration of the $\text{NH}_3 (g)$ at equilibrium.

(A) $9.0 \times 10^{-2} M$

(B) $3.0 \times 10^{-1} M$

(C) $1.0 \times 10^2 M$

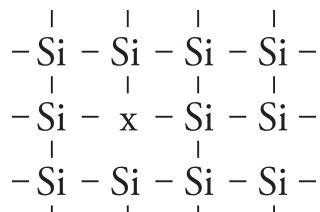
(D) $9.0 \times 10^2 M$

54. If some additional $\text{NH}_3 (g)$ is injected into the flask, which of the following options correctly predicts the reaction shift and what would happen to the concentration of NH_4HS ?

	Shift	$[\text{NH}_4\text{HS}]$
(A)	Left	Increase
(B)	Left	No Change
(C)	Right	Decrease

(D)	None	No Change
-----	------	-----------

55. If the temperature of the flask were to be increased to 313 K, which of the following identifies what would happen to the amount of NH_4HS present at equilibrium and correctly explains why?
- (A) Increase because both reaction rates have been permanently increased
- (B) Decrease because the value for K has increased
- (C) Decrease because the reaction is becoming more disordered
- (D) Remain the same because it is a solid and unaffected by reaction shifts
56. If the reaction were to be repeated at 298 K using the same initial amount of NH_4HS , but in a 500 mL flask, how would that affect the concentration of NH_3 at equilibrium compared to the trial in the 250 mL flask?
- (A) It would be unchanged.
- (B) It would decrease by a factor of four.
- (C) It would decrease by a factor of two.
- (D) It would increase by a factor of two.



57. A semiconductor with a silicon lattice is doped by the addition of another element, symbolized by the letter X in the above diagram. Which of the following statements best describes the

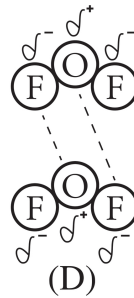
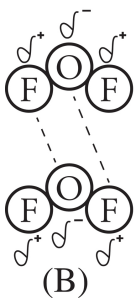
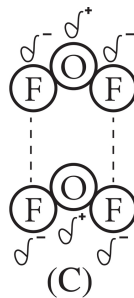
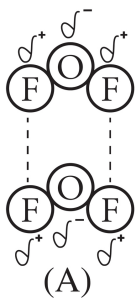
possible identity of the element as well as the type of doping that is occurring?

- (A) Boron, which has less valence electrons than silicon and creates positive “holes” in the lattice
- (B) Phosphorus, which has extra valence electrons and creates a mobile negative charge in the lattice
- (C) Aluminum, which has metalloid properties that lend to increased conductivity when added to a semiconductor
- (D) Carbon, which has an equal number of valence electrons as silicon and strengthens the conducting lattice

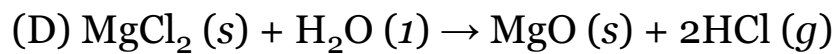
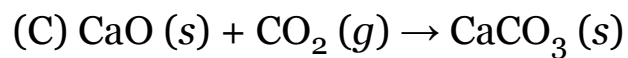
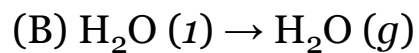
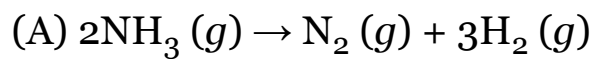
58. What is the oxidation state on the sulfur atom in the compound S_2Cl_2 ?

- (A) -2
- (B) -1
- (C) 0
- (D) $+1$

59. Which of the following diagrams correctly demonstrates the permanent dipole attractions between two molecules of oxygen difluoride, OF_2 ? Note that the dashed lines are representative of attractive intermolecular forces.



60. Which of the following reactions would have the most negative value for ΔS° ?



END OF SECTION I

Section II

INFORMATION IN THE TABLE BELOW AND ON THE FOLLOWING PAGES MAY BE USEFUL IN ANSWERING THE QUESTIONS IN THIS SECTION OF THE EXAMINATION.

DO NOT DETACH FROM BOOK.

PERIODIC TABLE OF THE ELEMENTS

1 H 1.008																	2 He 4.00
3 Li 6.94	4 Be 9.01											5 B 10.81	6 C 12.01	7 N 14.01	8 O 16.00	9 F 19.00	10 Ne 20.18
11 Na 22.99	12 Mg 24.30											13 Al 26.98	14 Si 28.09	15 P 30.97	16 S 32.06	17 Cl 35.45	18 Ar 39.95
19 K 39.10	20 Ca 40.08	21 Sc 44.96	22 Ti 47.90	23 V 50.94	24 Cr 52.00	25 Mn 54.94	26 Fe 55.85	27 Co 58.93	28 Ni 58.69	29 Cu 63.55	30 Zn 65.39	31 Ga 69.72	32 Ge 72.59	33 As 74.92	34 Se 78.96	35 Br 79.90	36 Kr 83.80
37 Rb 85.47	38 Sr 87.62	39 Y 88.91	40 Zr 91.22	41 Nb 92.91	42 Mo 95.94	43 Tc (98)	44 Ru 101.1	45 Rh 102.91	46 Pd 106.42	47 Ag 107.87	48 Cd 112.41	49 In 114.82	50 Sn 118.71	51 Sb 121.75	52 Te 127.60	53 I 126.91	54 Xe 131.29
55 Cs 132.91	56 Ba 137.33	*57 *La 138.91	72 Hf 178.49	73 Ta 180.95	74 W 183.85	75 Re 186.21	76 Os 190.2	77 Ir 192.2	78 Pt 195.08	79 Au 196.97	80 Hg 200.59	81 Tl 204.38	82 Pb 207.2	83 Bi 208.98	84 Po (209)	85 At (210)	86 Rn (222)
87 Fr (223)	88 Ra 226.02	†89 †Ac 227.03	104 Rf (261)	105 Db (262)	106 Sg (266)	107 Bh (264)	108 Hs (277)	109 Mt (268)	110 Ds (271)	111 Rg (272)							

*Lanthanide Series	58 Ce 140.12	59 Pr 140.91	60 Nd 144.24	61 Pm (145)	62 Sm 150.4	63 Eu 151.97	64 Gd 157.25	65 Tb 158.93	66 Dy 162.50	67 Ho 164.93	68 Er 167.26	69 Tm 168.93	70 Yb 173.04	71 Lu 174.97
†Actinide Series	90 Th 232.04	91 Pa 231.04	92 U 238.03	93 Np (237)	94 Pu (244)	95 Am (243)	96 Cm (247)	97 Bk (247)	98 Cf (251)	99 Es (252)	100 Fm (257)	101 Md (258)	102 No (259)	103 Lr (262)

ADVANCED PLACEMENT CHEMISTRY EQUATIONS AND CONSTANTS

Throughout the test the following symbols have the definitions specified unless otherwise noted.

L, mL = liter(s), milliliter(s)
 g = gram(s)
 nm = nanometer(s)
 atm = atmosphere(s)

mm Hg = millimeters of mercury
 J, kJ = joule(s), kilojoule(s)
 V = volt(s)
 mol = mole(s)

ATOMIC STRUCTURE

$$E = h\nu$$

$$c = \lambda\nu$$

E = energy
 ν = frequency
 λ = wavelength

Planck's constant, $h = 6.626 \times 10^{-34}$ J s

Speed of light, $c = 2.998 \times 10^8$ m s⁻¹

Avogadro's number = 6.022×10^{23} mol⁻¹

Electron charge, $e = -1.602 \times 10^{-19}$ coulomb

EQUILIBRIUM

$$K_c = \frac{[C]^c[D]^d}{[A]^a[B]^b}, \text{ where } aA + bB \rightleftharpoons cC + dD$$

$$K_p = \frac{(P_C)^c(P_D)^d}{(P_A)^a(P_B)^b}$$

$$K_a = \frac{[H^+][A^-]}{[HA]}$$

$$K_b = \frac{[OH^-][HB^+]}{[B]}$$

$$K_w = [H^+][OH^-] = 1.0 \times 10^{-14} \text{ at } 25^\circ\text{C}$$

$$= K_a \times K_b$$

$$\text{pH} = -\log[H^+], \text{ pOH} = -\log[OH^-]$$

$$14 = \text{pH} + \text{pOH}$$

$$\text{pH} = \text{p}K_a + \log \frac{[A^-]}{[HA]}$$

$$\text{p}K_a = -\log K_a, \text{ p}K_b = -\log K_b$$

Equilibrium Constants

K_c (molar concentrations)

K_p (gas pressures)

K_a (weak acid)

K_b (weak base)

K_w (water)

KINETICS

$$\ln[A]_t - \ln[A]_0 = -kt$$

$$\frac{1}{[A]_t} - \frac{1}{[A]_0} = kt$$

$$t_{1/2} = \frac{0.693}{k}$$

k = rate constant

t = time

$t_{1/2}$ = half-life

GASES, LIQUIDS, AND SOLUTIONS

$$PV = nRT$$

$$P_A = P_{\text{total}} \times X_A, \text{ where } X_A = \frac{\text{moles A}}{\text{total moles}}$$

$$P_{\text{total}} = P_A + P_B + P_C + \dots$$

$$n = \frac{m}{M}$$

$$K = ^\circ\text{C} + 273$$

$$D = \frac{m}{V}$$

$$KE \text{ per molecule} = \frac{1}{2}mv^2$$

Molarity, M = moles of solute per liter of solution

$$A = abc$$

P = pressure

V = volume

T = temperature

n = number of moles

m = mass

M = molar mass

D = density

KE = kinetic energy

v = velocity

A = absorbance

a = molar absorptivity

b = path length

c = concentration

Gas constant, $R = 8.314 \text{ J mol}^{-1} \text{ K}^{-1}$

$= 0.08206 \text{ L atm mol}^{-1} \text{ K}^{-1}$

$= 62.36 \text{ L torr mol}^{-1} \text{ K}^{-1}$

$1 \text{ atm} = 760 \text{ mm Hg}$

$= 760 \text{ torr}$

STP = 273.15 K and 1.0 atm

Ideal gas at STP = 22.4 L mol^{-1}

THERMOCHEMISTRY/ ELECTROCHEMISTRY

$$q = mc\Delta T$$

$$\Delta S^\circ = \sum S^\circ \text{ products} - \sum S^\circ \text{ reactants}$$

$$\Delta H^\circ = \sum \Delta H_f^\circ \text{ products} - \sum \Delta H_f^\circ \text{ reactants}$$

$$\Delta G^\circ = \sum \Delta G_f^\circ \text{ products} - \sum \Delta G_f^\circ \text{ reactants}$$

$$\Delta G^\circ = \Delta H^\circ - T\Delta S^\circ$$

$$= -RT \ln K$$

$$= -nFE^\circ$$

$$I = \frac{q}{t}$$

q = heat

m = mass

c = specific heat capacity

T = temperature

S° = standard entropy

H° = standard enthalpy

G° = standard free energy

n = number of moles

E° = standard reduction potential

I = current (amperes)

q = charge (coulombs)

t = time (seconds)

Faraday's constant, $F = 96,485 \text{ coulombs per mole of electrons}$

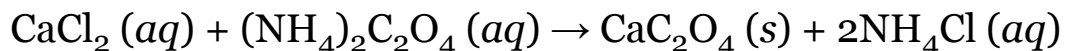
$$1 \text{ volt} = \frac{1 \text{ joule}}{1 \text{ coulomb}}$$

CHEMISTRY
Section II
7 Questions
(Total time—105 minutes)

YOU MAY USE YOUR CALCULATOR FOR THIS SECTION

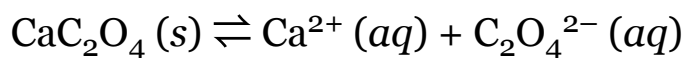
Directions: Questions 1-3 are long free-response questions that require about 23 minutes each to answer and are worth 10 points each. Questions 4-7 are short free-response questions that require about 9 minutes each to answer and are worth 4 points each.

Write your response in the space provided following each question. Examples and equations may be included in your responses where appropriate. For calculations, clearly show the method used and the steps involved in arriving at your answers. You must show your work to receive credit for your answer. Pay attention to significant figures.



1. Most vitamin tablets consist of not only the active ingredient, but also other ingredients that are there to bind the vitamin together or coat it for easier swallowing. A calcium tablet with a mass of 1.49 g is crushed, and then dissolved in some concentrated HCl, creating aqueous calcium chloride. The tablet solution is then mixed with excess ammonium oxalate in a basic environment, causing calcium oxalate to precipitate.
 - (a) Describe a method to fully separate the precipitate from the solution and accurately measure its mass.
 - (b) The final mass of the CaC_2O_4 is determined to be 1.64 g.
 - (i) How many moles of calcium are present in the precipitate?
 - (ii) What is the mass percent of calcium in the original vitamin tablet?

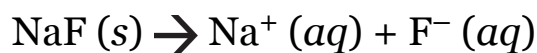
- (c) CaC_2O_4 has a K_{sp} value of 1.9×10^{-9} and dissociates in solution as follows.



- (i) What is the concentration of Ca^{2+} ions and $\text{C}_2\text{O}_4^{2-}$ ions in a saturated solution of CaC_2O_4 ?
- (ii) If some of the water evaporated from a saturated solution of CaC_2O_4 , how would that effect the concentration of the ions you determined in part (c)(i)? Justify your answer.

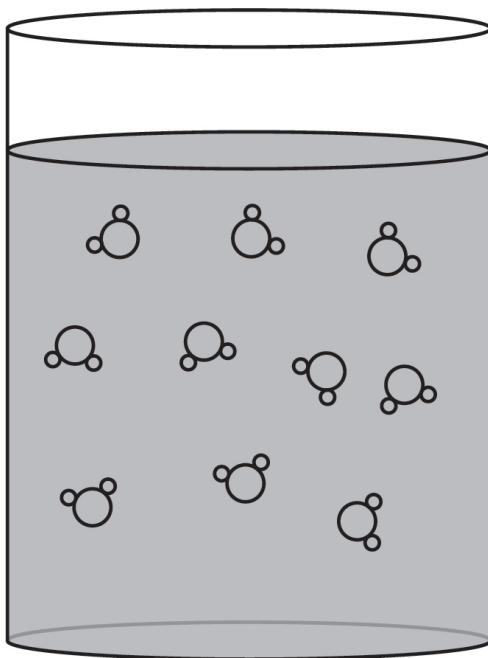
The results do not match the accepted value of calcium in the tablet provided by the manufacturer. Upon reviewing the reaction, it was determined that the final product was not, in fact, pure calcium oxalate, but was instead a hydrate of calcium oxalate.

- (d) Would not accounting for the water of hydration cause the calcium value in the tablet to be artificially high or artificially low? Why?
- (e) A 5.00 g sample of the calcium oxalate hydrate is heated several times, resulting in a final mass for the anhydrous salt of 4.38 g.
- (i) How many moles of water were present in the sample?
- (ii) What is the hydration coefficient?



2. Sodium fluoride is the active ingredient in toothpaste that helps prevent tooth decay. One of the important aspects of sodium fluoride is that it is completely soluble in water, and will dissociate fully.

- (a) What are the forces that exist between sodium fluoride and water molecules that allow for this dissociation to occur?
- (b) Which ion has the larger size- Na^+ or F^- ? Justify your answer.
- (c) The below beaker is filled with water molecules. In the beaker, correctly draw the location of the Na^+ and F^- ions in a dissolved solution of sodium fluoride. Make sure each ion is labeled clearly.



- (d) 5.00 g of NaF is dissolved into 50.0 mL of water, and the temperature of the water decreases by 0.52°C . The specific heat and density of solution are identical to that of pure water; $4.18 \text{ J/g}^\circ\text{C}$ and 1.0 g/mL , respectively.
- (i) How much energy is lost by the solution during the dissolution process?
- (ii) What is the enthalpy of solution, in kJ/mol , for the dissolution of sodium fluoride?

- (iii) For the dissolution of sodium fluoride, which value has a greater magnitude—the lattice energy of sodium fluoride, or the hydration energy between sodium fluoride and water? How do you know?
- (e) How would you expect the magnitude of the hydration energy for sodium fluoride compare to that of potassium chloride, KCl? Justify your answer.



3. Lactic acid is an acid found in many dairy-related products.

- (a) Write the equilibrium constant expression, K_a , for lactic acid.
- (b) (i) The pH of an 0.50 M solution of lactic acid is found to be 2.08. Calculate the percent dissociation of the 0.50 M lactic acid solution.
- (ii) If the 0.50 M solution from part (i) were to be diluted, how would that affect the percent dissociation, if at all? Justify your answer.
- (c) Lactic acid is the active ingredient in sour cream which makes it taste sour. A sour cream sample with a volume of 500. mL is titrated, and the lactic acid present in it is found to have concentration of 0.113 M . If the sour cream sample has a density of 1.0 g/mL, what is the percent by mass of lactic acid present in the sour cream?
- (d) To control the pH level of lactic acid containing substances, sodium lactate, $\text{NaC}_3\text{H}_5\text{O}_3$, is often used.
- (i) How many moles of sodium lactate would need to be added to 100. mL of 0.10 M lactic acid to create a buffer with a pH of 4.0?

- (ii) Adding a small amount of which of the following chemicals to a solution of lactic acid would also create a buffer solution? Justify your answer using a chemical reaction.

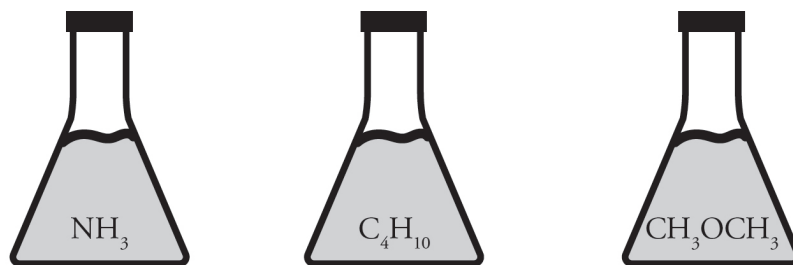


4. Many foods are dyed various colors to make them more aesthetically pleasing. The dyes are often complex organic compounds, including one that is commercially called Brilliant Blue FCF, which can be abbreviated as BB. When BB reacts with the active ingredient in bleach, the hypochlorite ion (ClO^-), the color will fade over time. Thus, the rate of reaction can be determined by using a colorimeter to measure the rate at which the BB fades.

To determine the reaction order for both BB and ClO^- in this reaction, three trials are run at 25°C and the rate of reaction is measured in each.

Trial	[BB] (M)	[ClO^-] (M)	Rate (M/s)
1	1.5×10^{-3}	1.5×10^{-3}	3.3×10^{-3}
2	2.0×10^{-3}	1.5×10^{-3}	4.4×10^{-3}
3	3.0×10^{-3}	2.5×10^{-3}	1.1×10^{-2}

- (a) What is the reaction order with respect to:
- (i) BB
 - (ii) ClO^-
- (b) Calculate the rate constant for the reaction between BB and ClO^- at 25°C . Include units.

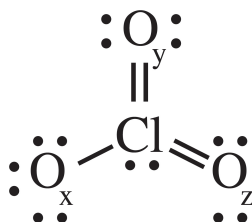


$$P_{\text{NH}_3} = 7,600 \text{ mmHg}$$

$$P_{\text{C}_4\text{H}_{10}} = 2,200 \text{ mmHg}$$

$$P_{\text{CH}_3\text{OCH}_3} = 200 \text{ mmHg}$$

5. Three different liquids are placed in three sealed flasks of identical volume as shown above. All of the flasks are held at a constant temperature of 25°C , and the vapor pressure of each gas is listed below the flask.
- Which liquid has the strongest intermolecular forces? Justify your answer on a particular level.
 - If the temperature of each of the flasks were to be increased, what effect would that have on the vapor pressure, if any? Why?
 - Are the gases present in each flask least likely to behave ideally at very low or very high temperatures? Explain your reasoning.
 - Which substance would show the greatest deviation from ideal behaviors under the conditions you chose in (c)(i)? Justify your answer.
6. Use the below diagram of the chlorate ion, ClO_3^- , to answer the following questions as needed.



- (a) What is the oxidation state on the chlorine atom in the chlorate ion? Show any necessary calculations.
- (b) Calculate the formal charge on each atom in the chlorate ion. Note that each oxygen atom should be clearly labeled to match the subscripts in the diagram.
- (c) Is the Cl-O_x bond shorter than, longer than, or the same length as the Cl-O_y bond? Justify your answer.
- (d) Chlorine atoms can expand their octet, but oxygen atoms cannot. Why?
7. The Br-Br bond in a bromine molecule has a bond energy of 193 kJ/mol.
- (a) (i) How much energy, in Joules, is required to break the bond in a single Br molecule?
- (ii) What wavelength of light is necessary to break a single Br-Br bond?
- (b) Is the wavelength of light necessary to break a F-F bond shorter than, longer than, or the same as the wavelength of light necessary to break the Br-Br bond? Justify your answer.

STOP

END OF EXAM

Practice Test 3: Answers and Explanations

PRACTICE TEST 3: MULTIPLE-CHOICE ANSWER KEY

1. D	21. A	41. B
2. B	22. A	42. A
3. A	23. C	43. B
4. D	24. A	44. D
5. D	25. B	45. C
6. C	26. C	46. A
7. A	27. B	47. A
8. A	28. B	48. D
9. A	29. A	49. B
10. D	30. A	50. B
11. B	31. C	51. C
12. C	32. D	52. C
13. C	33. A	53. B
14. D	34. C	54. B
15. A	35. B	55. B
16. A	36. A	56. A
17. C	37. A	57. A
18. C	38. C	58. D
19. D	39. D	59. D
20. B	40. C	60. C

SECTION I—MULTIPLE-CHOICE ANSWERS AND EXPLANATIONS

1. **D**

HOCl has the larger K_a , which means it dissociates more and produces more H^+ ions than HOBr. The reason for this is chlorine's electronegativity attracts the electrons bonded between O and H, which weakens that bond and makes the H more likely to dissociate.

2. **B**

NaOH has a molar mass of 40 g/mol. The oxygen present in the NaOH has a molar mass of 16 g/mol, and so makes up $16/40 = 40\%$ of the molar mass of NaOH. Oxygen would also represent 40% of the mass of the 10.0 g NaOH sample, which is 4.00 g.

3. **A**

Remember that Avogadro's Law states that one mole of ANY gas at STP takes up 22.4 L of Space. With that in mind:

$$3.5 \text{ g Li} \times \frac{1 \text{ mol Li}}{7.0 \text{ g Li}} \times \frac{1 \text{ mol H}_2}{2 \text{ mol Li}} \times \frac{22.4 \text{ L}}{1 \text{ mol H}_2} = 5.60 \text{ L}$$

4. **D**

The amount of heat gained by the water is equal to the amount of heat lost by the metal. This is mathematically represented by $q_{\text{water}} = -q_{\text{metal}}$. Using the calorimetry equation,

$$m_w c_w \Delta T_w = -m_m c_m \Delta T_m$$

$$(100)(4.2)(2) = -(10) c_m (-70)$$

$$840 = 700 c_m$$

$$c_m = 1.2 \text{ J/g}^\circ\text{C}$$

Note that even without a calculator, it should be pretty easy to see that c_m will be greater than 1 in the above equation, and in this case only one answer option is greater than one.

5. **D**

In an equilibrium constant expression, it is products over reactants and neither solids nor liquids are included due to their unchanging concentration.

6. **C**

Silver has the higher SRP and would be the cathode. To calculate cell potential, subtract the reduction potential of the anode from the reduction potential of the cathode.

$$0.80 \text{ V} - (-0.44 \text{ V}) = 1.24 \text{ V}$$

7. **A**

Electrons must be balanced in a full reaction, and to do that you must multiple the Ag half reaction by two before combining it with the Fe half reaction. The Fe half reaction will flip and become an oxidation because it has the lower SRP.

8. **A**

For metallic M ions with a charge of +Y:

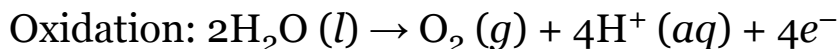
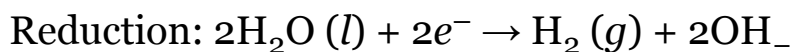
Current $\times$ time = Charge (in C)

$$\times \frac{1 \text{ mol } e^{-}}{96500 \text{ C}} \times \frac{1 \text{ mol M}}{Y \text{ mol } e^{-}} \times \frac{\text{Molar Mass of M}}{1 \text{ mol M}} = \text{mass M}$$

Given that silver has both the lowest charge and the highest molar mass, both factors lead to more silver metal being plated out.

9. **A**

When a current is run through water, it will both reduce into hydrogen gas and oxidize into oxygen gas using the given reactions.



The reduction reaction has to be multiplied by two to balance to charge, and the final reaction becomes: $2\text{H}_2\text{O} (l) \rightarrow 2\text{H}_2 (g) + \text{O}_2 (g)$ (H^{+} and OH^{-} combine to form H_2O , some of which cancels). Thus, twice as much H_2 is produced, meaning the H_2 is in the tube above electrode X, making electrode X the cathode as well.

10. **D**

When a solid metal is placed into a solution containing aqueous cations of another metal, the SRP of the metal in solution must exceed that of the solid metal for the solution to be able to take electrons from the solid and cause a reaction to occur. The only choice in which this does not occur (i.e., in which the SRP of the metal $>$ SRP of the solution) is (D).

11. **B**

Aluminum's electron configuration represented by the PES is $1s^2 2s^2 2p^6 3s^2 3p^1$. The $2p$ peak is the third-lowest energy level that an electron can be found in, and that peak is found at about 150 eV.

12. **C**

To form an ion, aluminum loses three valence electrons. Those would come from the $3s$ and $3p$ energy levels, removing those peaks from the spectrum.

13. **C**

The energy from the incoming radiation has to exceed the binding energy to cause electron ejection. 200 eV is not enough energy to eject a $1s$ or $2s$ electron. Comparing the $2p$ and $3p$ orbitals, the binding energy for a $2p$ electron is greater, meaning there will be less kinetic energy left post-ejection.

14. **D**

An amp is a coulomb per second.

$$4.0 \text{ C/s} = \frac{q}{120 \text{ s}} \quad q = 480 \text{ C}$$

15. **A**

The configuration of an oxygen atom is $1s^2 2s^2 2p^4$. The $2p$ sub-level has three orbitals, and one electron must fill each of them before any of them pair up.

16. **A**

All of the carbon will end up in the carbon dioxide, and all of the hydrogen will end up in the water. 44.0 g of CO_2 is equal to one mole, and with a 1:1 ratio that means there is one mole of carbon too. 18.0 g of H_2O is equal to one mole of H_2O , but as each mole of H_2O has two moles of hydrogen, that means there are four moles of hydrogen present. The carbon to hydrogen ratio in the original sample is thus 1:4, which matches the ratio in option (A).

17. **C**

The maximum absorbance of the copper ion occurs at approximately 640 nm. When studying a copper solution of unknown concentration, comparing it to a Beer's Law plot constructed at an absorbance of 640 nm would create the greatest possible absorbance range and give the best results.

18. **C**

The tin starts at an oxidation state of 0 in its elemental state, and ends up at an oxidation state of +2 in Sn^{2+} . This requires a loss of electrons, thus tin is oxidized. To determine the oxidation state of the chromium in dichromate, realize that oxygen is -2 in the compound and the total of the oxidation states equals the charge on the ion (-2). $2(\text{Cr}) + 7(-2) = -2$. $\text{Cr} = +6$. The chromium in Cr^{3+} ends up at an oxidation state of +3, meaning it gained electrons and thus was reduced.

19. **D**

Looking at the oxidation half-reaction, tin goes from 0 to +2, and does it three times in the balanced equation. That means 6 moles of electrons are transferred. You get the

same answer when looking at the reduction half-reaction with chromium going from +6 to +3 twice.

20. **B**

$$0.033 M = \frac{n}{0.100 L} \quad n = 0.0033 \text{ mol}$$

$$\text{K}_2\text{Cr}_2\text{O}_7 \times \frac{3 \text{ mol Sn}}{1 \text{ mol K}_2\text{Cr}_2\text{O}_7} = 0.010 \text{ mol Sn}$$

$$0.010 \text{ mol Sn} \times \frac{118.71 \text{ g}}{1 \text{ mol Sn}} = 1.2 \text{ g Sn}$$

21. **A**

When a salt dissociates in solution, whether or not it is endothermic or exothermic depends on the sum of the hydration energy and the lattice energy. Hydration energy is always negative, and lattice energy is always positive. Thus, if the value for the enthalpy of solution is becoming more positive, the (positive) lattice energy is increasing at a faster rate than the (negative) hydration energy.

22. **A**

An increase in the reverse reaction rate is a shift to the left. In an aqueous equilibrium, diluting the system will cause a shift to the side with more ions. In this equilibrium, there are three ions on the left and only one on the right, so adding water shifts the system left.

23. **C**

$$Q = \frac{[\text{Cu}^{2+}]}{[\text{Ag}^+]}$$

K for any galvanic cell is very large, and if Q ever reaches K, the cell will be at equilibrium and have zero voltage. For this cell, increasing $[Ag^+]$ causes Q to decrease, taking it further from K, which means it is also further from equilibrium and further from zero voltage.

24. **A**

$$0.50 \text{ M NaOH} = \frac{n}{0.020 \text{ L}} \quad n = 0.010 \text{ mol}$$
$$\text{NaOH} \times \frac{1 \text{ mol HN}_3}{1 \text{ mol NaOH}} = \frac{0.010 \text{ mol HN}_3}{0.025 \text{ L}} = 0.40 \text{ M}$$

25. **B**

At equivalence (20.0 mL), all of the HN_3 would be converted to N^{3-} . At half-equivalence, only half of the HN_3 would be converted to N^{3-} , meaning the concentration of both would be equal.

26. **C**

Indicators change color at their pK_a value. The best indicator for a titration is one that changes color at the equivalence point of the titration. For this titration that occurs at about a pH of 9, meaning the thymol blue is the best choice.

27. **B**

The amount the pH changes at the very beginning of the titration is dependent on the K_a of the weak acid. The lower the K_a , the weaker the acid is and the more the addition of NaOH will change the pH of the solution. Note that (D) is wrong because a weaker acid will have a stronger conjugate

base, meaning that at equivalence the solution will be more basic (higher pH).

28. **B**

All three ions are isoelectric, meaning they have the same number of electrons and the same electron configuration. However, the nucleus of the aluminum ion has the most protons, and can pull the electrons in closer. This makes them harder to remove and gives the aluminum ion the greatest ionization energy.

29. **A**

The negative dipoles on the water molecules are located on the oxygen end of the molecule. These dipoles would be pointing toward the sodium cation (and the positive hydrogen dipoles would point toward the chloride anion). The sodium ion is also smaller than the chloride ion, leading to the correct answer.

30. **A**

The equation you need here is $d = P(MM)/RT$, where MM is the molar mass of the gas (in this case, 4.0 g/mol). At STP, the pressure is 1.0 atm and the temperature is 273 K, while the ideal gas constant is always 0.0821 atm*L/mol*K.

31. **C**

If you multiply the bottom reaction by four and flip it, it turns into $2\text{O}_2(g) + 4\text{H}_2(g) \rightarrow 4\text{H}_2\text{O}(l)$. Combining that with the unchanged top two reactions yields the target reaction.

32. **D**

When flipping the initial reaction, take the reciprocal of the equilibrium constant. $1/(5.0 \times 10^{-3}) = 2.0 \times 10^2$. Then, when the coefficients are doubled, the reaction coefficient must be squared. $(2.0 \times 10^2)^2 = 4.0 \times 10^4$.

33. **A**

The substance starts at a solid, and begins to melt at 50°C . This is known because the temperature stops changing, which is always true during a phase change. While it is melting, the solid and liquid phases will exist simultaneously.

34. **C**

In the gas phase, which occurs at point D, the intermolecular forces present are very weak due to the great distance between the gas molecules as well as their high velocities.

35. **B**

The amount of heat needed to change the substance from a liquid to a gas (heat of vaporization) exceeds the amount of heat needed to change it from a solid to a liquid (heat of fusion). This is based on the length of the horizontal lines representing each phase change.

36. **A**

The solid to liquid phase change (the first section of unchanging temperature) requires 200 J of energy to complete. A 1.0 sample of the substance would require half as much heat to melt, so 100 J.

37. **A**

The substance is in the liquid phase starting at 50°C and ending at 150°C. During that time, 400 J of heat is added. Using $q = mc\Delta T$: $400 \text{ J} = (2.0 \text{ g})c(100^\circ\text{C})$ $c = 2.0 \text{ J/g}^\circ\text{C}$

38. **C**

When a weak acid is diluted, the reverse reaction becomes more hindered because the H^+ and F^- ions do not collide as often due to the overwhelming number of reactants surrounding them. This has the effect of increasing the percent dissociation as the H^+ and F^- ions are more likely to remain in solution. Note that (D) is wrong because a weak acid cannot be dissociated 100%; if it were, that would make it a strong acid.

39. **D**

Voltage is a function of the reaction quotient for the battery. Given that $Q = [\text{Zn}^{2+}]/[\text{Cu}^{2+}]$, in both batteries the value for Q would be the same, making their voltages the same. However, battery Y has both more metal ions in solution and in the electrodes, meaning the battery could operate for longer.

40. **C**

$\text{C}_6\text{H}_{12}\text{O}_6$ is a covalent substance, so that would have the lowest melting point. Of the three ionic substances, Mg^{2+} and O^{2-} are both smaller than and have a greater magnitude of charge than the ions present in the other salts. These factors would lead to greater lattice energy and a higher melting point.

41. **B**

If all gases are at the same temperature, they have the same amount of kinetic energy. For this to be true, the gas with the lowest molar mass must have the fastest particles.

42. **A**

With number of moles and temperature constant, pressure and volume have an inverse relationship defined using $P_1V_1 = P_2V_2$. Treating each gas separately:

$$\text{Kr: } (1.0)(1.0) = P_2(3.0) \quad P_2 = 0.33 \text{ atm}$$

$$\text{H}_2\text{O: } (2.0)(1.0) = P_2(3.0) \quad P_2 = 0.67 \text{ atm}$$

$$\text{CO}_2: (3.0)(1.0) = P_2(3.0) \quad P_2 = 1.0 \text{ atm}$$

$$P_T = 0.33 + 0.67 + 1.0 = 2.0 \text{ atm}$$

43. **B**

Pressure and temperature have a direct relationship which is linear in nature. As temperature increases, pressure will increase at a proportional rate.

44. **D**

$\text{C}_2\text{H}_5\text{OH}$ and PF_3 are covalent substances, which means they are poor conductors of electricity. Of the two ionic substances, NH_4Cl breaks down into two ions (NH_4^+ and Cl^-) while Na_2CO_3 breaks down into three (2 Na^+ and CO_3^{2-}). More ions in solution = greater conductivity.

45. **C**

The boiling point is based on the strength of the intermolecular forces between the halogen molecules. All

are non-polar, so the only type of IMF present is London dispersion forces. The strength of the LDFs is based in large part on how many electrons there are present, and more electrons create stronger temporary dipoles, leading to stronger overall IMFs.

46. **A**

When determining the enthalpy of reaction using bond enthalpies, bonds broken in the reactants are assigned a positive value and bonds formed in the products are assigned a negative value prior to summing them all up. Remember to account for both the number of bonds inside a molecule as well as how many of that molecule there is in the balanced equation when calculating (for instance, there are 4 H-O bonds in 2 H₂O molecules).

$$4(\text{C-H}) + (\text{C}=\text{C}) + 3(\text{O}=\text{O}) - 4(\text{C}=\text{O}) - 4(\text{H-O}) = \Delta H_{\text{rxn}}^{\circ}$$

$$4(410) + 720 + 3(500) - 4(800) - 4(470) = -1220 \text{ kJ/mol}$$

47. **A**

$$K_p = \frac{(\text{PNO})^2(\text{PO}_2)}{(\text{PNO}_2)^2} \quad K_p = \frac{(3 \times 10^{-1})^2(2 \times 10^{-1})}{(6 \times 10^{-1})^2}$$

$$K_p = \frac{18 \times 10^{-3}}{36 \times 10^{-2}} \quad K_p = 0.5 \times 10^{-1} = .05$$

48. **D**

The bond angle is based on the number of charge clouds (either a lone pair or any type of bond) around the central atom in the bond. There are only three charge clouds around the carbon in the C-C=O bond, making the bond angle about 120°. There are four charge clouds that are the

central atoms of the H-C-H and H-N-H bonds, meaning the bond angle is approximately 109.5° . However, one of those charge clouds on the H-N-H bond is a lone pair, which exerts slightly more repulsive force than a bonded pair, shrinking the bond angle.

49. **B**

All single bonds are sigma bonds, but any additional shared pairs of electrons in double or triple bonds are pi bonds. There's only one double bond present, and so there is only one pi bond (since the first bonded pair is still a sigma bond).

50. **B**

Hydrogen bonding is the strongest type of intermolecular force, and occurs when hydrogen is bonded to either nitrogen, oxygen, or fluorine. In acetamide, there are two hydrogens bonded directly to the nitrogen, creating hydrogen bonding.

51. **C**

The ClO_2^- ion is the conjugate base of HClO_2 , and for any conjugate pair $K_a \times K_b = 1.0 \times 10^{-14}$. So, the K_b for ClO_2^- is 1.0×10^{-12} . From there, solve for the weak-base equilibrium.



$$K_b = \frac{[\text{HClO}_2][\text{OH}^-]}{[\text{ClO}_2^-]} \quad 1.0 \times 10^{-12} = \frac{x^2}{1.0 \text{ M}}$$

$$x = 1.0 \times 10^{-6} = [\text{OH}^-]$$

$$-\log[\text{OH}^-] = \text{pOH} \quad -\log(1.0 \times 10^{-6}) = \text{pOH} \quad \text{pOH} = 6$$

$$\text{pH} + \text{pOH} = 14 \quad \text{pH} + 6 = 14 \quad \text{pH} = 8$$

52. **C**

Between trial 1 and 2, the concentration of SO_2 remains the same but the concentration of O_3 changes. However, the rate does not change, which means the O_3 concentration has no effect on rate, making O_3 zero order. Between trials 1 and 3, the concentration of O_3 remains the same while the concentration of SO_2 doubles. The rate increases by a factor of four, making the reaction second order with respect to SO_2 .

53. **B**

$$K_c = \frac{[\text{H}_2\text{S}][\text{NH}_3]}{[\text{NH}_3]} \quad 9.0 \times 10^{-2} = (x)(x) \quad x = 3.0 \times 10^{-1} =$$

54. **B**

Adding the NH_3 would cause the equilibrium to shift left in order to relieve the stress. However, the NH_4HS is a solid, and solids have a set concentration which cannot be changed. That is also why adding or removing a solid (or liquid) from an equilibrium system does not cause a shift.

55. **B**

This reaction is endothermic, meaning an increase in temperature causes a permanent shift to the right. That would increase the value for K , meaning there are more H_2S and NH_3 present once equilibrium re-establishes. Notice that while the concentration of NH_4HS is a solid and cannot change, the amount can still change. In this case, creating more products means there is less reactant left.

56. **A**

As long as the temperature remains constant, so does the value for K . If K is constant, so are the concentrations of all the products and reactants at equilibrium.

57. **A**

Element X only forms three bonds with the surrounding silicon atoms, meaning it only has three available valence electrons to form bonds. That is consistent with boron, and adding boron into a silicon lattice is called p-doping, which involves creating positive areas for any mobile electrons to be attracted to.

58. **D**

The most electronegative element in a compound will have an oxidation state that is equal to the most common charge of its corresponding ion. In this case, chlorine is more electronegative than sulfur, and would thus have an oxidation state of -1 . The entire molecular must have a combined oxidation state of zero, so to balance the two chlorines each sulfur must be $+1$.

59. **D**

Fluorine is more electronegative than oxygen, so fluorine would have a negative dipole and oxygen would have a positive dipole in OF_2 . The attraction between two molecules would be of the negative (F) dipole on one molecule to the positive (O) dipole on the other.

60. **C**

A negative value for ΔS° means the entropy is decreasing, so the system is becoming more ordered. The phase change

from a gas to a solid in (C) demonstrates this, while all other options show entropy increasing.

SECTION II—FREE-RESPONSE ANSWERS AND EXPLANATIONS

1. Most vitamin tablets consist of not only the active ingredient, but also other ingredients that are there to bind the vitamin together or coat it for easier swallowing. A calcium tablet with a mass of 1.49 g is crushed, and then dissolved in some concentrated HCl, creating aqueous calcium chloride. The tablet solution is then mixed with excess ammonium oxalate in a basic environment, causing calcium oxalate to precipitate.

- (a) Describe a method to fully separate the precipitate from the solution and accurately measure its mass.

First, the solution containing the precipitate needs to be poured into a funnel that has been prepared with a piece of filter paper. The solution should be filtered as many times as needed until the filtrate is clear. The filter paper and precipitate should then be fully dried prior to a final massing.

- (b) The final mass of the CaC_2O_4 is determined to be 1.64 g.

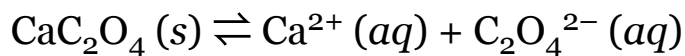
- (i) How many moles of calcium are present in the precipitate?

$$1.64 \text{ g CaC}_2\text{O}_4 \times \frac{1 \text{ mol CaC}_2\text{O}_4}{128.10 \text{ g}} \times \frac{1 \text{ mol Ca}}{1 \text{ mol CaC}_2\text{O}_4} = 0.0128 \text{ mol Ca}$$

- (ii) What is the mass percent of calcium in the original vitamin tablet?

$$\frac{0.0128 \text{ mol Ca} \times \frac{40.08 \text{ g}}{1 \text{ mol Ca}}}{\frac{0.513 \text{ g Ca}}{1.49 \text{ g}}} \times 100\% = 34.4\% \text{ Ca}$$

- (c) CaC_2O_4 has a K_{sp} value of 1.9×10^{-9} and dissociates in solution as follows.



- (i) What is the concentration of Ca^{2+} ions and $\text{C}_2\text{O}_4^{2-}$ ions in a saturated solution of CaC_2O_4 ?

$K_{\text{sp}} = [\text{Ca}^{2+}][\text{C}_2\text{O}_4^{2-}]$ The ions exist in a 1:1 ratio, so both unknowns can be replaced with x .

$$K_{\text{sp}} = (x)(x) \quad 1.9 \times 10^{-9} = x^2 \quad x = [\text{Ca}^{2+}] = [\text{C}_2\text{O}_4^{2-}] = 4.4 \times 10^{-5} \text{ M}$$

- (ii) If some of the water evaporated from a saturated solution of CaC_2O_4 , how would that effect the concentration of the ions you determined in part (c)(i)? Justify your answer.

The concentration of both ions will remain the same. For a saturated solution, the ions are already present at their maximum possible concentration values. If some water evaporates, some of the ions that were suspended between the evaporate water molecules will precipitate out, leaving the total concentration unchanged.

The results do not match the accepted value of calcium in the tablet provided by the manufacturer. Upon reviewing the reaction, it was determined that the final product was

not, in fact, pure calcium oxalate, but was instead a hydrate of calcium oxalate.

- (d) Would not accounting for the water of hydration cause the calcium value in the tablet to be artificially high or artificially low? Why?

If the calcium oxalate is a hydrate, that means there will be some waters of hydration attached to the crystalline lattice. This will cause the mass to increase, meaning the mass that was used in the calculations is artificially low, making the moles of calcium and the calcium percentage in the tablet artificially high.

- (e) A 5.00 g sample of the calcium oxalate hydrate is heated several times, resulting in a final mass for the anhydrous salt of 4.38 g.

- (i) How many moles of water were present in the sample?

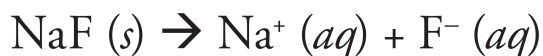
$$\begin{aligned} 5.00 \text{ g} - 4.38 \text{ g} &= 0.62 \text{ g H}_2\text{O} \times \\ \frac{1 \text{ mol H}_2\text{O}}{18.01 \text{ g H}_2\text{O}} &= 0.034 \text{ mol H}_2\text{O} \end{aligned}$$

- (ii) What is the hydration coefficient?

To determine the hydration coefficient, we have to compare the moles of water to the moles of the anhydrous salt.

$$4.38 \text{ g CaC}_2\text{O}_4 \times \frac{1 \text{ mol CaC}_2\text{O}_4}{128.10 \text{ g}} = 0.034 \text{ mol CaC}_2\text{O}_4$$

The moles of water and moles of salt exist in a 1:1 ratio, therefore the hydration coefficient is 1.



2. Sodium fluoride is the active ingredient in toothpaste that helps prevent tooth decay. One of the important aspects of sodium fluoride is that it is completely soluble in water, and will dissociate fully.

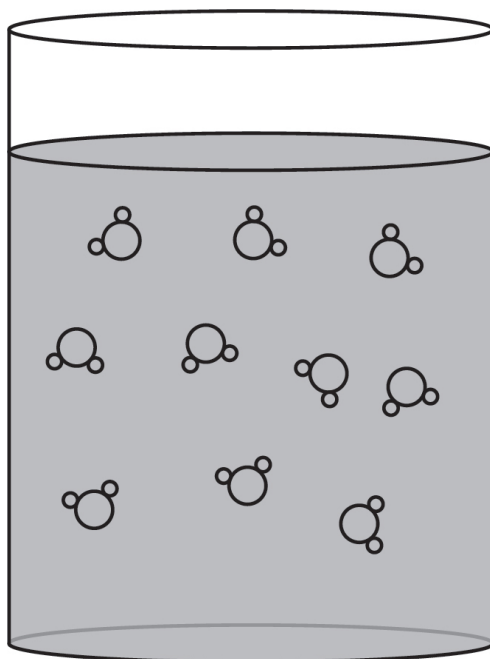
- (a) What are the forces that exist between sodium fluoride and water molecules that allow for this dissociation to occur?

An ionic salt dissolves when the attraction of its ions to the dipoles in the water molecules exceeds the attraction of the ions to each other. In this case, the positive sodium cations will be attracted to the negative water dipoles, and the negative fluoride ions will be attracted to the positive water dipoles.

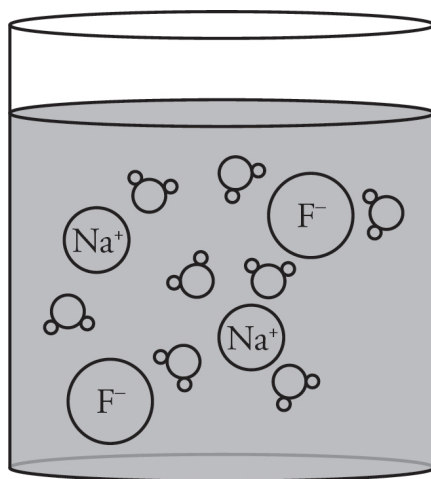
- (b) Which ion has the larger size— Na^+ or F^- ? Justify your answer.

F^- is larger. Both ions have the same number of electrons⁻ 10. When looking at species that have the same number of electrons, the one with more protons (in this case, Na^+) will be smaller, because the nuclear charge is greater and can pull those electrons in closer.

- (c) The below beaker is filled with water molecules. In the beaker, correctly draw the location of the Na^+ and F^- ions in a dissolved solution of sodium fluoride. Make sure each ion is labeled clearly.



The Na^+ ions must be between the negative dipoles, which in water are on the oxygen atoms. The F^- ions must be between the positive water dipoles located on the hydrogen atoms.



- (d) 5.00 g of NaF is dissolved into 50.0 mL of water, and the temperature of the water decreases by 0.52°C . The specific heat and density of solution are identical to that of pure water; $4.18 \text{ J/g}^\circ\text{C}$ and 1.0 g/mL , respectively.

- (i) How much energy is lost by the water during the dissolution process?

$q = mc\Delta T$ is the equation needed here. Note that the mass is the mass of the water plus the mass of the dissolved salt!

$$q = (55.0 \text{ g})(4.18 \text{ J/g}^\circ\text{C})(0.52^\circ\text{C})$$

$$q = -120. \text{ J}$$

- (ii) What is the enthalpy of solution, in kJ/mol, for the dissolution of sodium fluoride?

If the water lost 120. J of energy, that means the dissolution process required 120. J of energy. The sign flips to indicate the energy gain.

$$\Delta H = \frac{q}{n} \quad q = 120. \text{ J}$$

$$n = 5.00 \text{ g NaF} \times \frac{1 \text{ mol NaF}}{42.0 \text{ g NaF}} = 0.119 \text{ mol NaF}$$

$$\Delta H = 120. \text{ J} / 0.119 \text{ mol} = 1010 \text{ J/mol} = 1.01 \text{ kJ/mol}$$

- (iii) For the dissolution of sodium fluoride, which value has a greater magnitude—the lattice energy of sodium fluoride, or the hydration energy between sodium fluoride and water? How do you know?

Lattice energy describes the amount of energy necessary to break the bonds in an ionic substance. Breaking bonds requires energy, so lattice energy is a positive value.

Hydration energy is the amount of energy released when the dipoles of water molecules form attractions with the charged ions in the salt. Forming attractions releases energy, so hydration energy is always a negative value.

The total enthalpy of solution is the sum of the lattice and hydration energy values for a particular salt. In this case, the enthalpy of solution is positive, meaning the magnitude of the lattice energy exceeds that of the hydration energy.

- (e) How would you expect the magnitude of the hydration energy for sodium fluoride compare to that of potassium chloride, KCl? Justify your answer.

Hydration energy is based on Coulomb's Law. In this case, the charges of all the ions are the same—+1 on the cations and −1 on the anions. Looking at their sizes, however, Na^+ is smaller than K^+ and F^- is smaller than Cl^- . As Coulombic energy is inversely proportional with size, the substance with the smaller ions would have a greater Coulombic energy, and thus the greater hydration energy. So, the magnitude of NaF's hydration energy exceeds that of KCl.



3. Lactic acid is an acid found in many dairy-related products.

- (a) Write the equilibrium constant expression, K_a , for lactic acid

$$K_a = \frac{[\text{H}^+][\text{CH}_3\text{H}_5\text{O}_3^-]}{[\text{HC}_3\text{H}_5\text{O}_3]}$$

- (b) (i) The pH of an 0.50 M solution of lactic acid is found to be 2.08. Calculate the percent dissociation of the 0.50 M lactic acid solution.

$$-\log [\text{H}^+] = 2.08 \quad [\text{H}^+] = 10^{-2.08} \quad [\text{H}^+] = 8.3 \times 10^{-3} M$$

$$\% \text{ dissociation} = \frac{[\text{H}^+]}{[\text{HC}_3\text{H}_5\text{O}_3]} = \frac{8.3 \times 10^{-3} \text{ M}}{0.50 \text{ M}} \times 100 \% = 1.66\%$$

- (ii) If the 0.50 *M* solution from part (i) were to be diluted, how would that affect the percent dissociation, if at all? Justify your answer.

The percent dissociation would increase. The more dilute the solution is, the less likely the reverse reaction is to occur. This means a larger percentage of ions will stay dissociated.

- (c) Lactic acid is the active ingredient in sour cream which makes it taste sour. A sour cream sample with a volume of 500. mL is titrated, and the lactic acid present in it is found to have concentration of 0.113 *M*. If the sour cream sample has a density of 1.0 g/mL, what is the percent by mass of lactic acid present in the sour cream?

$$\begin{aligned} 0.113 \text{ M} &= \frac{n}{0.500 \text{ L}} & n &= 0.0565 \text{ mol HCH}_3\text{H}_5\text{O}_3 \\ &\times \frac{69.08 \text{ g}}{1 \text{ mol HCH}_3\text{H}_5\text{O}_3} & &= 3.90 \text{ g HCH}_3\text{H}_5\text{O}_3 \\ &\frac{3.09 \text{ g}}{500. \text{ g}} \times 100 \% & &= 0.78\% \text{ lactic acid} \end{aligned}$$

- (d) To control the pH level of lactic acid containing substances, sodium lactate, NaC₃H₅O₃, is often used.

- (i) How many moles of sodium lactate would need to be added to 100. mL of 0.10 *M* lactic acid to create a buffer with a pH of 4.0?

Using Henderson-Haselbalch:

$$\text{pH} = \text{p}K_{\text{a}} + \log \frac{[\text{CH}_3\text{H}_5\text{O}_3^-]}{[\text{HC}_3\text{H}_5\text{O}_3]}$$

$$4.0 = -\log (1.38 \times 10^{-4}) + \log \frac{[\text{CH}_3\text{H}_5\text{O}_3^-]}{0.10 \text{ M}}$$

$$4.0 = 3.86 + \log \frac{[\text{CH}_3\text{H}_5\text{O}_3^-]}{0.10 \text{ M}}$$

$$0.14 = \log \frac{[\text{CH}_3\text{H}_5\text{O}_3^-]}{0.10 \text{ M}}$$

$$0.014 = \log [\text{CH}_3\text{H}_5\text{O}_3^-]$$

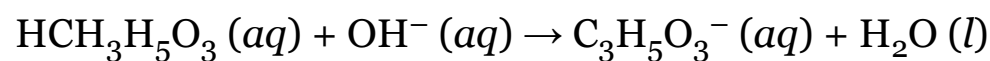
$$[\text{CH}_3\text{H}_5\text{O}_3^-] = 1.03 \text{ M}$$

$$1.03 \text{ M} = \frac{n}{0.100 \text{ L}}$$

$$n = 0.103 \text{ mol NaC}_3\text{H}_5\text{O}_3$$

- (ii) Adding a small amount of which of the following chemicals to a solution of lactic acid would also create a buffer solution? Justify your answer using a chemical reaction.

NaOH. Adding a strong base to a weak acid will cause the creation of conjugate base ions as follows:



Those conjugate base ions will create a buffer with the remaining weak acid.

HCl NaOH KCl

4. Many foods are dyed various colors to make them more aesthetically pleasing. The dyes are often complex organic compounds, including one that is commercially called Brilliant Blue FCF, which can be abbreviated as BB. When BB reacts with the active ingredient in bleach, the hypochlorite ion (ClO^-), the color will fade over time. Thus, the rate of reaction can be determined by using a colorimeter to measure the rate at which the BB fades.

To determine the reaction order for both BB and ClO^- in this reaction, three trials are run at 25°C and the rate of reaction is measured in each.

Trial	[BB] (M)	[ClO^-] (M)	Rate (M/s)
1	1.5×10^{-3}	1.5×10^{-3}	3.3×10^{-3}
2	2.0×10^{-3}	1.5×10^{-3}	4.4×10^{-3}
3	3.0×10^{-3}	2.5×10^{-3}	1.1×10^{-2}

(a) What is the reaction order with respect to:

(i) BB

To calculate the order with respect to BB, look at the first two trials. The concentration of the ClO^- stays constant, so any change in rate is due to the change in the initial concentration of the BB.

$$\frac{3.2 \times 10^{-3} = k(1.5 \times 10^{-3})^m(1.5 \times 10^{-3})^n}{4.4 \times 10^{-3} = k(2.0 \times 10^{-3})^m(1.5 \times 10^{-3})^n}$$

$$0.74 = 0.75^m \quad m = 1$$

The reaction is first order with respect to BB.

(ii) ClO^-

The concentration of BB changes with every trial, but because we know the order, we can still determine the order with respect to ClO^- by inputting the BB order into our equation. Using trials 2 and 3:

$$\frac{4.4 \times 10^{-3} = k(2.0 \times 10^{-3})^1(1.5 \times 10^{-3})^n}{1.1 \times 10^{-2} = k(3.0 \times 10^{-3})^1(2.5 \times 10^{-3})^n}$$

$$0.40 = (0.67)(0.60)^n \quad 0.60 = 0.60^n \quad n = 1$$

The reaction is first order with respect to ClO^- .

(b) Calculate the rate constant for the reaction between BB and ClO^- at 25°C .

Data from any trial can be plugged into the rate law, along with the determined orders, to figure out the rate constant. The determined rate law is $\text{Rate} = k[\text{BB}][\text{ClO}^-]$, and using data from trial 2 the magnitude of the rate constant can be determined.

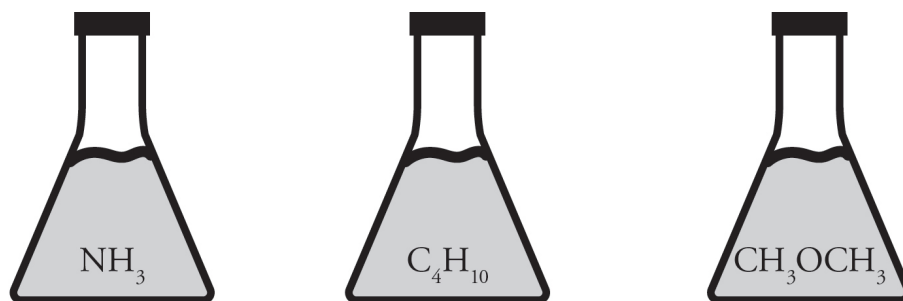
$$4.4 \times 10^{-3} = k(2.0 \times 10^{-3})(1.5 \times 10^{-3})$$

$$k = 1500$$

Using dimensional analysis, we can see that unitwise:

$$\text{M/s} = k(\text{M})(\text{M})$$

The units must be $\text{M}^{-1}\text{s}^{-1}$, so the final answer is $1500 \text{ M}^{-1}\text{s}^{-1}$.



$$P_{\text{NH}_3} = 7,600 \text{ mmHg}$$

$$P_{\text{C}_4\text{H}_{10}} = 2,200 \text{ mmHg}$$

$$P_{\text{CH}_3\text{OCH}_3} = 200 \text{ mmHg}$$

5. Three different liquids are placed in three sealed flasks of identical volume as shown above. All of the flasks are held at a constant temperature of 25°C , and the vapor pressure of each gas is listed below the flask.

- (a) Which liquid has the strongest intermolecular forces? Justify your answer on a particular level.

CH_3OCH_3 has the strongest IMFs, as shown by its low vapor pressure. The lower the vapor pressure, the less likely the molecules are to escape the liquid state, which requires a breaking of all IMFs. Stronger IMFs = less change of molecules escaping = lower vapor pressure.

- (b) If the temperature of each of the flasks were to be increased, what effect would that have on the vapor pressure, if any? Why?

The vapor pressure would increase. Adding energy to the liquids gives their molecules more kinetic energy, making them more likely to be able to overcome the IMFs holding them in the liquid state.

- (c) (i) Are the gases present in each flask least likely to behave ideally at very low or very high temperatures? Explain

your reasoning.

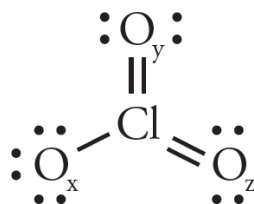
Deviations are more likely to occur at very low temperatures. This is because the gas molecules will have less energy, making them less likely to be able to overcome the IMFs between them.

In an ideal gas, IMFs are very weak, but the slower the gas molecules are moving the more likely the IMFs are to become significant enough to cause deviations from ideal behavior.

- (ii) Which substance would show the greatest deviation from ideal behaviors under the conditions you chose in (c)(i)? Justify your answer.

The gas with the strongest IMFs is most likely to deviate from ideal behavior, which in this case means the CH_3OCH_3 is most likely to behave non-ideally.

6. Use the below diagram of the chlorate ion, ClO_3^- , to answer the following questions as needed.



- (a) What is the oxidation state on the chlorine atom in the chlorate ion? Show any necessary calculations.

Oxygen has an oxidation state of negative two, and the total oxidation state on the chlorate must be equal to its charge of negative one. So: $\text{Cl} + 3(-2) = -1$ $\text{Cl} = +5$

- (b) Calculate the formal charge on each atom in the chlorate ion. Note that each oxygen atom should be clearly labeled

to match the subscripts in the diagram.

Formal charge is equal to the number of valence electrons minus the number of assigned electrons drawn in the structure. Lone pairs count as two assigned electrons, while each bond attached to an atom counts as one.

	Cl	O _x	O _y	O _z
Valence e^-	7	6	6	6
Assigned e^-	7	7	6	6
Formal Charge	0	-1	0	0

- (c) Is the Cl-O_x bond shorter than, longer than, or the same length as the Cl-O_y bond? Justify your answer.

They are the same length. The chlorate ion is drawn with resonance, meaning the bonds are neither single bonds nor double bonds. All bonds would be somewhere in length between true single and double bonds, and all bonds are identical.

- (d) Chlorine atoms can expand their octet, but oxygen atoms cannot. Why?

To expand an octet, there must be an empty “*d*” orbital available on the same principal energy level as the valence electrons for that atom. Chlorine’s valence electrons are in the third principal energy level, and a 3*d* orbital does exist. Oxygen, however, has its valence electrons in the second principal energy level, and there is no 2*d* orbital, meaning there is no place for the extra electrons to go.

7. The Br-Br bond in a bromine molecule has a bond energy of 193 kJ/mol.

- (a) (i) How much energy, in Joules, is required to break the bond in a single Br molecule?

$$\frac{193 \text{ kJ}}{\text{mol}} \times \frac{1 \text{ mol}}{6.02 \times 10^{23} \text{ molecules}} \times \frac{1000 \text{ J}}{1 \text{ kJ}} = 3.21 \times 10^{-19} \text{ J}$$

- (ii) What wavelength of light is necessary to break a single Br-Br bond?

$$E = h\nu \quad 3.21 \times 10^{-19} \text{ J} = 6.634 \times 10^{-34} \text{ Js } (\nu) \quad \nu = 4.83 \times 10^{14} \text{ s}^{-1}$$

$$c = \nu\lambda \quad 3.00 \times 10^8 \text{ m/s} = 4.83 \times 10^{14} \text{ s}^{-1} \lambda \quad \lambda = 6.21 \times 10^{-7} \text{ m} = 621 \text{ nm}$$

- (b) Is the wavelength of light necessary to break a F-F bond shorter than, longer than, or the same as the wavelength of light necessary to break the Br-Br bond? Justify your answer.

The bond in F-F is shorter than the bond in Br-Br because a fluorine atom is smaller than a bromine atom (and they are both single bonds). A shorter bond has greater Coulombic energy. Energy and wavelength are inversely proportional, so breaking a F-F bond would require light of a shorter wavelength than the Br-Br bond.

Practice Test 4

[Click here](#) to download and print the PDF version of this exercise.

AP[®] Chemistry Exam

SECTION I: Multiple-Choice Questions

DO NOT OPEN THIS BOOKLET UNTIL YOU ARE TOLD TO DO SO.

At a Glance

Total Time

1 hour and 30 minutes

Number of Questions

60

Percent of Total Grade

50%

Writing Instrument

Pencil required

Instructions

Section I of this examination contains 60 multiple-choice questions. Fill in only the ovals for numbers 1 through 60 on your answer sheet.

CALCULATORS MAY NOT BE USED IN THIS PART OF THE EXAMINATION.

Indicate all of your answers to the multiple-choice questions on the answer sheet. No credit will be given for anything written in this exam booklet, but you may use the booklet for notes or scratch work. After you have decided which of the suggested answers is best, completely fill in the corresponding oval on the answer sheet. Give only one answer to each question. If you change an answer, be sure that the previous mark is erased completely. Here is a sample question and answer.

Sample Question

Chicago is a

- (A) state
- (B) city
- (C) country
- (D) continent

Sample Answer



Use your time effectively, working as quickly as you can without losing accuracy. Do not spend too much time on any one question. Go on to other questions and come back to the ones you have not answered if you have time. It is not expected that everyone will know the answers to all the multiple-choice questions.

About Guessing

Many candidates wonder whether or not to guess the answers to questions about which they are not certain. Multiple-choice scores are based on the number of questions answered correctly. Points are not deducted for incorrect answers, and no points are awarded for unanswered questions. Because points are not deducted for incorrect answers, you are encouraged to answer all multiple-choice questions. On any questions you do not know the answer to, you should eliminate as many choices as you can, and then select the best answer among the remaining choices.

Section I

CHEMISTRY SECTION I

Time—1 hour and 30 minutes

INFORMATION IN THE TABLE BELOW AND ON THE FOLLOWING PAGES MAY BE USEFUL IN ANSWERING THE QUESTIONS IN THIS SECTION OF THE EXAMINATION.

DO NOT DETACH FROM BOOK.

PERIODIC TABLE OF THE ELEMENTS

PERIODIC TABLE OF THE ELEMENTS																	
1 H 1.008																2 He 4.00	
3 Li 6.94	4 Be 9.01											5 B 10.81	6 C 12.01	7 N 14.01	8 O 16.00	9 F 19.00	10 Ne 20.18
11 Na 22.99	12 Mg 24.30											13 Al 26.98	14 Si 28.09	15 P 30.97	16 S 32.06	17 Cl 35.45	18 Ar 39.95
19 K 39.10	20 Ca 40.08	21 Sc 44.96	22 Ti 47.90	23 V 50.94	24 Cr 52.00	25 Mn 54.94	26 Fe 55.85	27 Co 58.93	28 Ni 58.69	29 Cu 63.55	30 Zn 65.39	31 Ga 69.72	32 Ge 72.59	33 As 74.92	34 Se 78.96	35 Br 79.90	36 Kr 83.80
37 Rb 85.47	38 Sr 87.62	39 Y 88.91	40 Zr 91.22	41 Nb 92.91	42 Mo 95.94	43 Tc (98)	44 Ru 101.1	45 Rh 102.91	46 Pd 106.42	47 Ag 107.87	48 Cd 112.41	49 In 114.82	50 Sn 118.71	51 Sb 121.75	52 Te 127.60	53 I 126.91	54 Xe 131.29
55 Cs 132.91	56 Ba 137.33	57 *La 138.91	72 Hf 178.49	73 Ta 180.95	74 W 183.85	75 Re 186.21	76 Os 190.2	77 Ir 192.2	78 Pt 195.08	79 Au 196.97	80 Hg 200.59	81 Tl 204.38	82 Pb 207.2	83 Bi 208.98	84 Po (209)	85 At (210)	86 Rn (222)
87 Fr (223)	88 Ra 226.02	89 †Ac 227.03	104 Rf (261)	105 Db (262)	106 Sg (266)	107 Bh (264)	108 Hs (277)	109 Mt (268)	110 Ds (271)	111 Rg (272)							

*Lanthanide Series

58 Ce 140.12	59 Pr 140.91	60 Nd 144.24	61 Pm (145)	62 Sm 150.4	63 Eu 151.97	64 Gd 157.25	65 Tb 158.93	66 Dy 162.50	67 Ho 164.93	68 Er 167.26	69 Tm 168.93	70 Yb 173.04	71 Lu 174.97
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†Actinide Series

90 Th 232.04	91 Pa 231.04	92 U 238.03	93 Np (237)	94 Pu (244)	95 Am (243)	96 Cm (247)	97 Bk (247)	98 Cf (251)	99 Es (252)	100 Fm (257)	101 Md (258)	102 No (259)	103 Lr (262)
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ADVANCED PLACEMENT CHEMISTRY EQUATIONS AND CONSTANTS

Throughout the test the following symbols have the definitions specified unless otherwise noted.

L, mL = liter(s), milliliter(s)
 g = gram(s)
 nm = nanometer(s)
 atm = atmosphere(s)

mm Hg = millimeters of mercury
 J, kJ = joule(s), kilojoule(s)
 V = volt(s)
 mol = mole(s)

ATOMIC STRUCTURE

$$E = h\nu$$

$$c = \lambda\nu$$

E = energy
 ν = frequency
 λ = wavelength

Planck's constant, $h = 6.626 \times 10^{-34}$ J s

Speed of light, $c = 2.998 \times 10^8$ m s⁻¹

Avogadro's number = 6.022×10^{23} mol⁻¹

Electron charge, $e = -1.602 \times 10^{-19}$ coulomb

EQUILIBRIUM

$$K_c = \frac{[C]^c[D]^d}{[A]^a[B]^b}, \text{ where } aA + bB \rightleftharpoons cC + dD$$

$$K_p = \frac{(P_C)^c(P_D)^d}{(P_A)^a(P_B)^b}$$

$$K_a = \frac{[H^+][A^-]}{[HA]}$$

$$K_b = \frac{[OH^-][HB^+]}{[B]}$$

$$K_w = [H^+][OH^-] = 1.0 \times 10^{-14} \text{ at } 25^\circ\text{C}$$

$$= K_a \times K_b$$

$$\text{pH} = -\log[H^+], \text{ pOH} = -\log[OH^-]$$

$$14 = \text{pH} + \text{pOH}$$

$$\text{pH} = \text{p}K_a + \log \frac{[A^-]}{[HA]}$$

$$\text{p}K_a = -\log K_a, \text{ p}K_b = -\log K_b$$

Equilibrium Constants

K_c (molar concentrations)

K_p (gas pressures)

K_a (weak acid)

K_b (weak base)

K_w (water)

KINETICS

$$\ln[A]_t - \ln[A]_0 = -kt$$

$$\frac{1}{[A]_t} - \frac{1}{[A]_0} = kt$$

$$t_{1/2} = \frac{0.693}{k}$$

k = rate constant

t = time

$t_{1/2}$ = half-life

GASES, LIQUIDS, AND SOLUTIONS

$$PV = nRT$$

$$P_A = P_{\text{total}} \times X_A, \text{ where } X_A = \frac{\text{moles A}}{\text{total moles}}$$

$$P_{\text{total}} = P_A + P_B + P_C + \dots$$

$$n = \frac{m}{M}$$

$$K = ^\circ\text{C} + 273$$

$$D = \frac{m}{V}$$

$$KE \text{ per molecule} = \frac{1}{2}mv^2$$

Molarity, M = moles of solute per liter of solution

$$A = abc$$

P = pressure

V = volume

T = temperature

n = number of moles

m = mass

M = molar mass

D = density

KE = kinetic energy

v = velocity

A = absorbance

a = molar absorptivity

b = path length

c = concentration

Gas constant, $R = 8.314 \text{ J mol}^{-1} \text{ K}^{-1}$

$= 0.08206 \text{ L atm mol}^{-1} \text{ K}^{-1}$

$= 62.36 \text{ L torr mol}^{-1} \text{ K}^{-1}$

$1 \text{ atm} = 760 \text{ mm Hg}$

$= 760 \text{ torr}$

STP = 273.15 K and 1.0 atm

Ideal gas at STP = 22.4 L mol^{-1}

THERMOCHEMISTRY/ ELECTROCHEMISTRY

$$q = mc\Delta T$$

$$\Delta S^\circ = \sum S^\circ \text{ products} - \sum S^\circ \text{ reactants}$$

$$\Delta H^\circ = \sum \Delta H_f^\circ \text{ products} - \sum \Delta H_f^\circ \text{ reactants}$$

$$\Delta G^\circ = \sum \Delta G_f^\circ \text{ products} - \sum \Delta G_f^\circ \text{ reactants}$$

$$\Delta G^\circ = \Delta H^\circ - T\Delta S^\circ$$

$$= -RT \ln K$$

$$= -nFE^\circ$$

$$I = \frac{q}{t}$$

q = heat

m = mass

c = specific heat capacity

T = temperature

S° = standard entropy

H° = standard enthalpy

G° = standard free energy

n = number of moles

E° = standard reduction potential

I = current (amperes)

q = charge (coulombs)

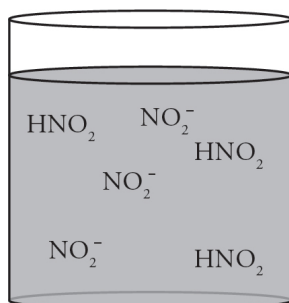
t = time (seconds)

Faraday's constant, $F = 96,485 \text{ coulombs per mole of electrons}$

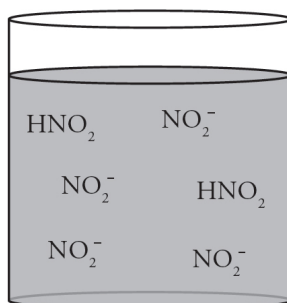
$$1 \text{ volt} = \frac{1 \text{ joule}}{1 \text{ coulomb}}$$



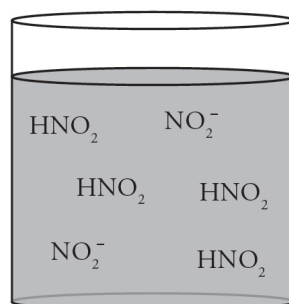
1. Which of the following is true about a beaker containing a saturated solution of silver carbonate?
 - (A) The molar solubility of Ag_2CO_3 is equal to the concentration of the silver ions.
 - (B) There would be no solid present at the bottom of the beaker.
 - (C) The concentration of the silver ions will be twice that of the carbonate ions.
 - (D) Adding more water would increase the solubility product constant.
2. Nitrous acid, HNO_2 , has a $\text{p}K_{\text{a}}$ value of 3.4. A solution of nitrous acid has some sodium nitrite, NaNO_2 , added to it, creating a buffer solution. If the pH of the buffer solution is found to be 3.4, which of the following particulate representations shows the correct ratio of acid and conjugate base molecules in the buffer?



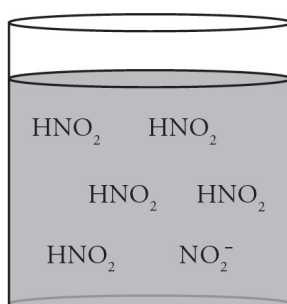
(A)



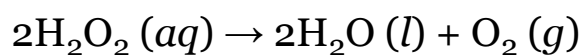
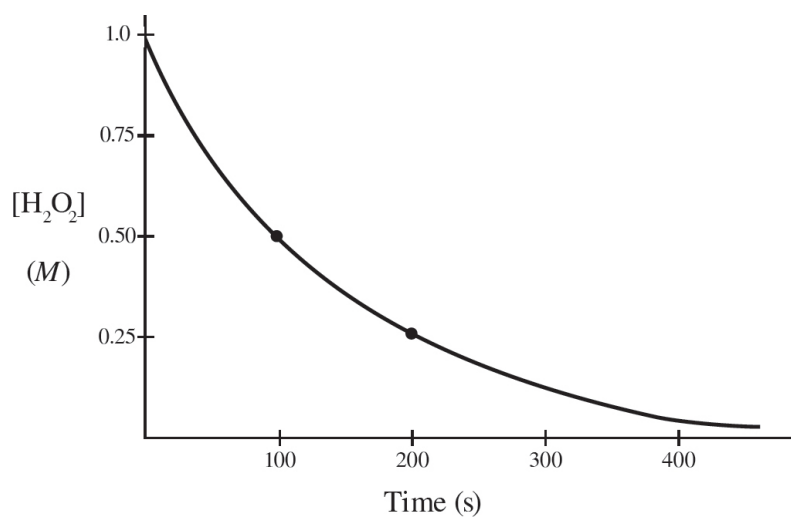
(C)



(B)

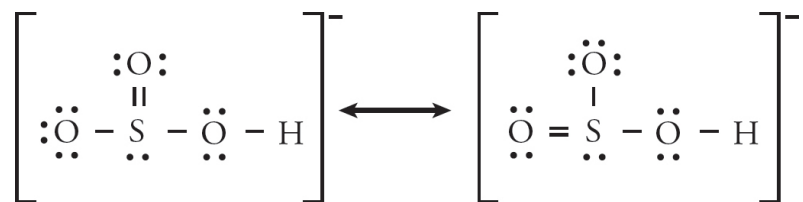
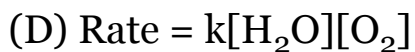
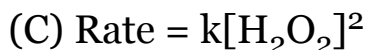
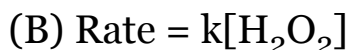


(D)



3. A sample of hydrogen peroxide, H_2O_2 , will decompose over time. The above graph charts the concentration of a H_2O_2 sample over time. What is the rate law for the overall reaction?

(A) Rate = k



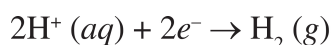
4. The Lewis diagram of the bisulfite ion, HSO_3^- , is drawn above. Which of the following descriptions correctly compares the bond lengths of the three sulfur-oxygen bonds found in bisulfite?

(A) All bonds are the same length.

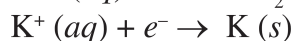
(B) Of the three bonds, there is one longer one and two shorter ones of identical length.

(C) Of the three bonds, there are two longer ones of identical length and one shorter one.

(D) All three bonds are different lengths.

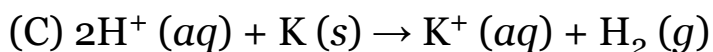
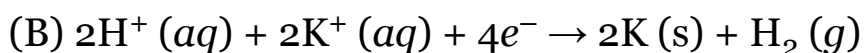
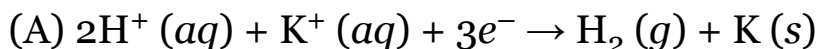


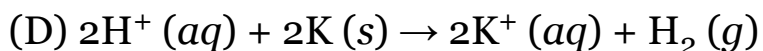
$$E^\circ = +0.00 \text{ V}$$



$$E^\circ = -2.92 \text{ V}$$

5. A piece of potassium metal is added to a solution of hydrochloric acid, and the metal starts to react. Which of the following is the correct net ionic equation for the reaction that is occurring?

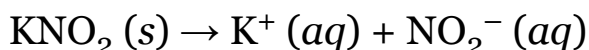




6. A solution of methanol, $CH_3OH (aq)$, has a vapor pressure of 13.2 kPa at room temperature. If the solution were to be heated, what would happen to the vapor pressure, and why?

- (A) Increase, because intramolecular attractions are breaking
(B) Increase, because the molecules in solution are moving faster
(C) Decrease, because stronger intermolecular attractions will form
(D) Remain unchanged, because the system is already at equilibrium

Use the following information to answer questions 7-10.



7. A student wants to determine the enthalpy of solution for potassium nitrite, KNO_2 (molar mass = 85.1 g/mol). The student takes a sample of KNO_2 and fully dissolves it into some water that is in a Styrofoam cup, gathering the following data. The density and specific heat of the final solution are identical to that of pure water, 1.0 g/mL and 4.2 J/g°C, respectively.

Mass KNO_2	8.50 g
Volume Water	91.50 mL
Initial Water Temperature	22.5°C
Final Water Temperature	19.5°C

What is the approximate enthalpy of solution for KNO_2 ?

- (A) -12.6 kJ/mol

(B) -1.26 kJ/mol

(C) 1.26 kJ/mol

(D) 12.6 kJ/mol

8. Which of the following statements is correct regarding the favorability of this process?

(A) The process is favored and driven by enthalpy and entropy.

(B) The process is favored and driven by enthalpy only.

(C) The process is favored and driven by entropy only.

(D) The process is not favored.

9. For the reaction to be endothermic, the magnitude of the attractive forces between the dissociated ions and the dipoles in the water molecules must be smaller which of the following?

(A) The intermolecular forces between the water molecules

(B) The attractive forces between the K^+ and NO_3^- ions in solution

(C) The covalent bond strength within the water molecules

(D) The strength of the ionic bonds in the KNO_3 (s) lattice

10. After the data is gathered, the student notices that not all of the KNO_2 sample dissolved. How would that affect the calculated enthalpy of solution?

(A) The calculated enthalpy value would be artificially low.

(B) The calculated enthalpy value would be artificially high.

(C) The calculated enthalpy value would be unchanged.

(D) The calculated enthalpy value would have the incorrect sign.



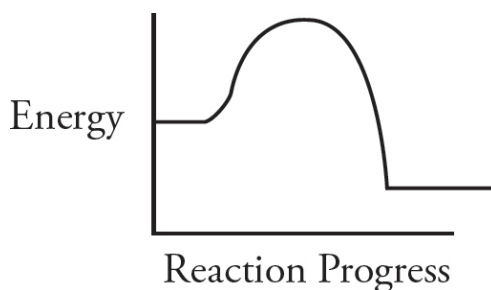
11. Strontium iodate is a slightly soluble salt that dissociates in water via the above equation. What concentration of Sr^{2+} ions would be necessary to precipitate $\text{Sr}(\text{IO}_3)_2$ in a solution that contains IO_3^- ions at a concentration of 0.010 *M*?

- (A) 0.0010 *M*
 (B) 0.010 *M*
 (C) 0.10 *M*
 (D) 1.0 *M*

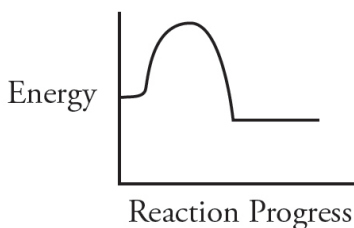
Acid	K_{a1}	K_{a2}	K_{a3}
Formic	1.8×10^{-4}	N/A	N/A
Sulfurous	1.5×10^{-2}	1.0×10^{-7}	N/A
Citric	8.4×10^{-4}	1.8×10^{-5}	4.0×10^{-6}

12. The K_{a} values for three different weak acids are given above. If the acid is polyprotic, all of the K_{a} values for that acid are given. If each acid were to be titrated with 0.10 *M* NaOH, which solution would have the highest pH at the endpoint of the titration when each acid has fully reacted?

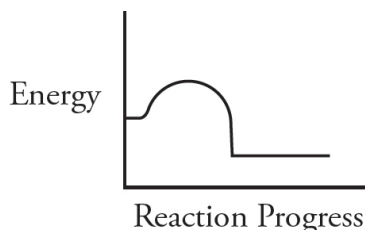
- (A) Formic
 (B) Sulfurous
 (C) Citric
 (D) All titrations would have identical pHs at the endpoint.



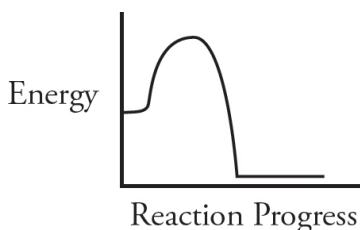
13. The above reaction coordinate shows the energy profile of an uncatalyzed reaction. If the reaction is repeated under identical collisions but a catalyst is added, which of the following coordinates is an accurate representation of what the energy profile of the catalyzed reaction could look like?



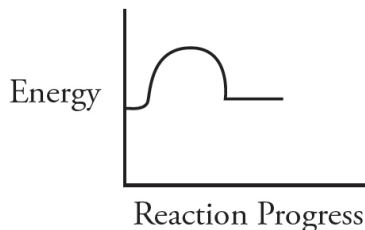
(A)



(C)



(B)



(D)

Use the following information to answer questions 14-18.

Name	Formula	K_a
Chlorous	HClO_2	1.0×10^{-2}
Carbonic	H_2CO_3	4.3×10^{-7}
Hypobromous	HBrO	2.0×10^{-9}

The K_a values for the first dissociation of three different weak acids are given in the above table.

14. A 0.10 M solution of which acid would have the highest concentration of H^+ ions?
- (A) $HClO_2$
 - (B) H_2CO_3
 - (C) $HBrO$
 - (D) $[H^+]$ would be equal in all solutions
15. Identify the conjugate base of hypobromous acid.
- (A) BrO^-
 - (B) H_2BrO^+
 - (C) H_3O^+
 - (D) OH^-
16. Which of the following solutions would create a buffer system with chlorous acid when added in an equimolar amount?
- (A) HCl
 - (B) $NaOH$
 - (C) $KClO_2$
 - (D) $HOCl$
17. Calculate the approximate pH of a 1.0 M solution of chlorous acid.
- (A) 1
 - (B) 2

(C) 3

(D) 4

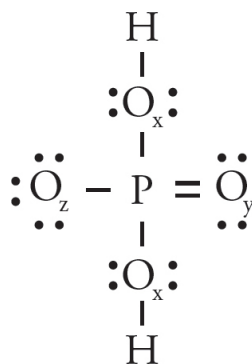
18. Carbonic acid is a polyprotic acid, and the second K_a value is 4.8×10^{-11} . Which of the following best explains why the K_a for the second dissociation is lower than that of the first?

(A) HCO_3^- is a more effective base than CO_3^{2-} .

(B) Acid dissociations are generally endothermic processes, and the first dissociation cools the reaction mixture, lowering the K_a value.

(C) Fewer water molecules are present to accept protons from HCO_3^- after the first dissociation has occurred.

(D) The presence of H^+ ions in solution after the first dissociation hinders the second dissociation.



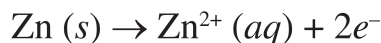
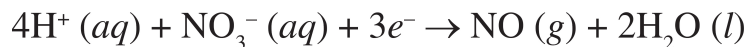
19. A Lewis diagram of the hydrogen phosphate ion is shown above. Based on this diagram, which of the atoms would have a negative formal charge?

(A) P

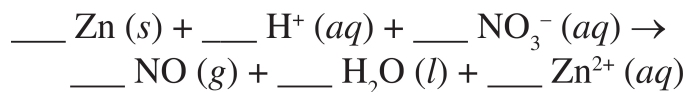
(B) O_x

(C) O_y

(D) O_z



20. Given the above half-reactions, what would the coefficient on the NO (g) be when the full reaction below is balanced?



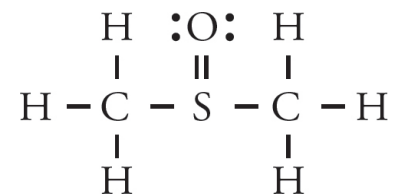
(A) 1

(B) 2

(C) 3

(D) 4

Use the following information to answer questions 21-23.



The Lewis diagram of dimethyl sulfoxide, commonly known as DMSO, is shown above.

21. What is the orbital hybridization found around the sulfur atom?

(A) sp

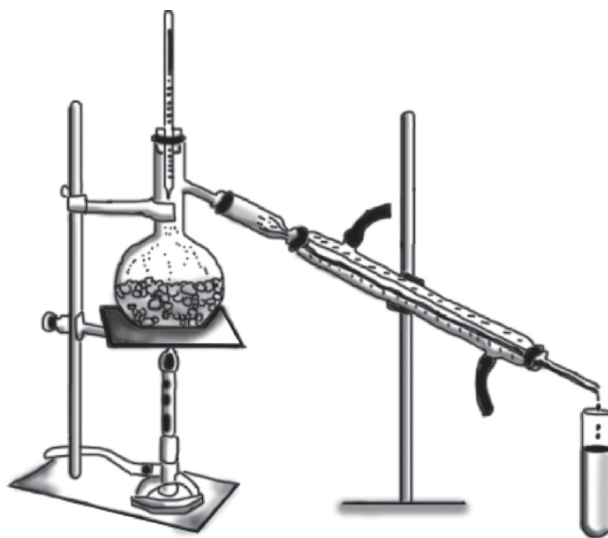
(B) sp^2

(C) sp^3

(D) sp^4

22. Which of the following bonds would have the longest length?

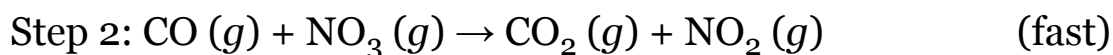
- (A) S-O
 - (B) S-C
 - (C) C-H
 - (D) All bonds would be equal in length
23. Where would the strongest dipoles on a molecule of DMSO be located?
- (A) Negative dipole on O, positive dipole on S
 - (B) Negative dipole on S, positive dipole on C
 - (C) Negative dipole on O, positive dipole on C
 - (D) The molecule is completely nonpolar.



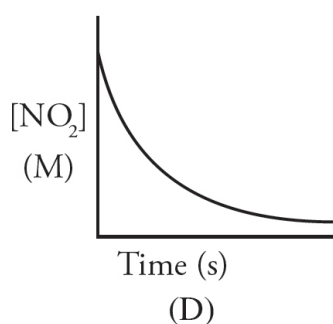
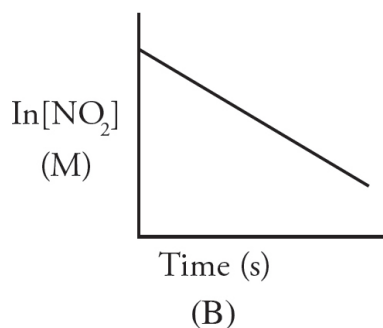
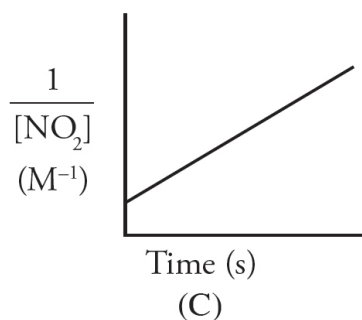
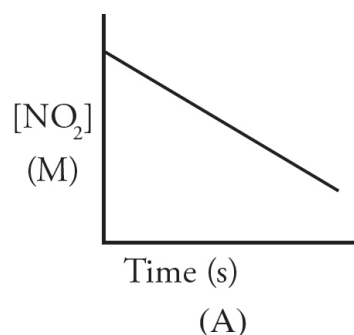
24. A student is attempting to use a distillation apparatus, shown above, to separate a mixture of propanol (boiling point: 97°C) and propionic acid (boiling point: 141°C). Which temperature should the apparatus be set at to obtain as pure a distillate of propanol as possible?
- (A) 85°C
 - (B) 105°C

(C) 125°C

(D) 145°C



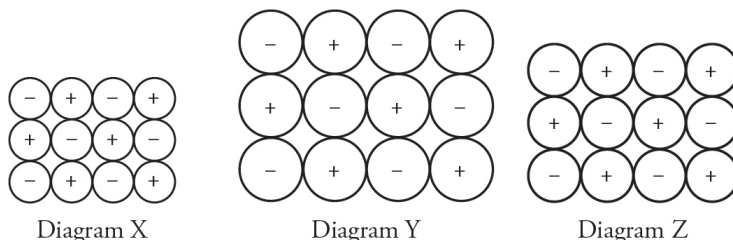
25. The above reaction mechanism shows the accepted elementary steps for a reaction. Which of the following graphs accurately represents the concentration of NO_2 versus time for the full reaction?



26. Which type of radiation would be best for studying the electron configuration of an individual atom?

- (A) Microwave
- (B) Infrared
- (C) Visible
- (D) Ultraviolet

Use the following information to answer questions 27-29.



The three diagrams above are particulate-level diagrams of three ionic solids: LiF, NaCl, and KBr. The diagrams are not organized in any particular order, and in each diagram the cations are marked with a + sign and the anions are marked with a – sign. All ions are drawn to scale.

27. Which diagram belongs to which ionic salt?

	Diagram X	Diagram Y	Diagram Z
(A)	LiF	NaCl	KBr
(B)	KBr	NaCl	LiF
(C)	KBr	LiF	NaCl
(D)	NaCl	KBr	LiF

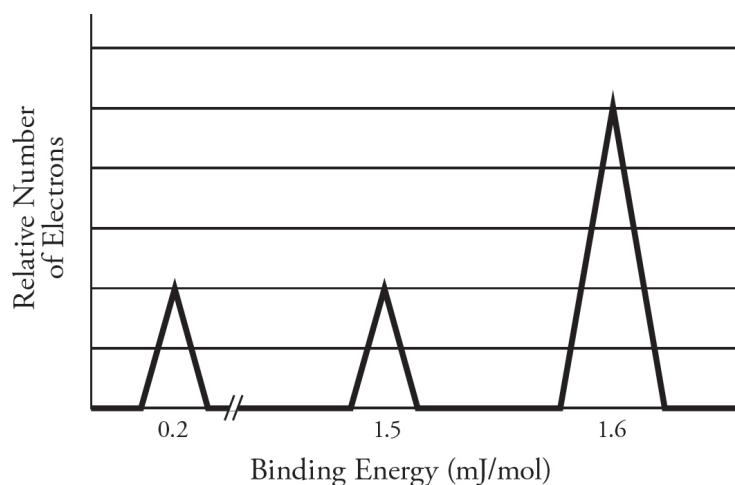
28. Which salt would have the highest melting point?

- (A) LiF
- (B) NaCl
- (C) KBr
- (D) The melting points of all three salts would be identical.

29. Which of the following ions would have a completed p-subshell in their valence level?

- (A) All the cations only (Li^+ , Na^+ , and K^+)

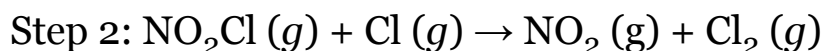
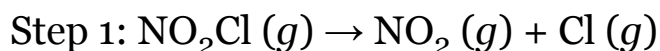
- (B) All the anions only (F^- , Cl^- , and Br^-)
- (C) Only K^+ and Br^-
- (D) All ions except for Li^+
30. Methanol, CH_3OH , is heated until it boils. During the boiling process, which of the following statements is true?
- (A) The speed of the molecules is constant.
- (B) The bonds within the molecules are breaking.
- (C) The intermolecular forces in the liquid are growing stronger.
- (D) The methanol is reacting with atmospheric gases.



31. The above photoelectron spectrum is for a neutral fluorine atom. Compared to that spectrum, the spectrum for a fluoride ion would have:
- (A) one more peak
- (B) one less peak
- (C) the same number of peaks, but the rightmost peak would be taller
- (D) the same number of peaks, but the leftmost peak would be taller

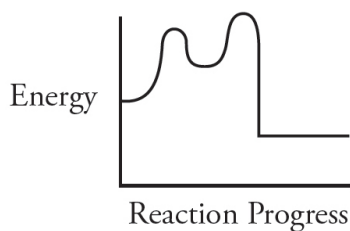
32. Determine the bond order for the B-F bond in a molecule of boron trifluoride, BF_3 .
- (A) 1
(B) 1.33
(C) 1.67
(D) 2
33. Copper (II) sulfate pentahydrate, $\text{CuSO}_4 \times 5 \text{H}_2\text{O}$, has a molar mass of 250 g/mol. How many grams of oxygen are present in a 25.0 g sample of $\text{CuSO}_4 \times 5 \text{H}_2\text{O}$?
- (A) 6.40 g
(B) 8.00 g
(C) 11.2 g
(C) 14.4 g
34. A sample of a noble gas at STP with a mass of 10.0 g is found to have a volume of 5.6 L. Which of the following gases correctly identifies the gas in the flask?
- (A) Helium
(B) Neon
(C) Argon
(D) Krypton

Use the following information to answer questions 35-38.

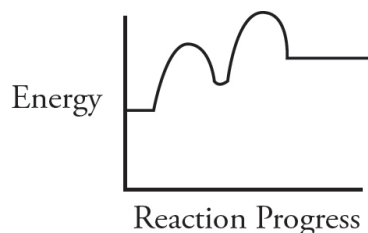


A proposed mechanism for a reaction is given above. The rate law for the overall reaction is known to be $\text{rate} = k[\text{NO}_2\text{Cl}]$.

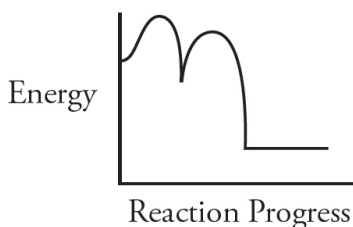
35. What is the molecularity of the slowest elementary step?
- (A) Unimolecular
 - (B) Bimolecular
 - (C) Trimolecular
 - (D) Tetramolecular
36. Which of the following could be the correct units on the rate constant, k ?
- (A) s^{-1}
 - (B) M^{-1}
 - (C) $\text{M}^{-1}\text{s}^{-1}$
 - (D) $\text{M}^{-1}\text{s}^{-2}$
37. Identify any catalysts and intermediates present in the proposed mechanism.
- (A) Catalyst: NO_2Cl Intermediate: Cl
 - (B) Catalyst: None Intermediate: Cl
 - (C) Catalyst: Cl Intermediate: None
 - (D) There are neither catalysts nor intermediates in the proposed mechanism.
38. The overall reaction is known to be exothermic. Which of the following energy profiles represents a potential energy pathway for the total reaction?



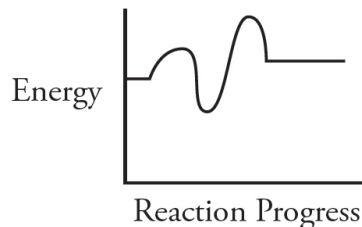
(A)



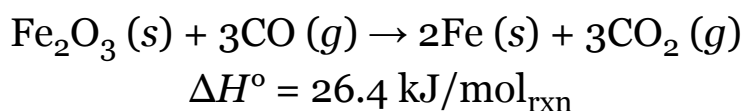
(C)



(B)



(D)



39. How much heat is released or absorbed when 2.80 g of CO reacts with excess Fe_2O_3 via the above reaction?

- (A) 2.63 kJ is released
- (B) 0.88 kJ is released
- (C) 2.63 kJ is absorbed
- (D) 0.88 kJ is absorbed



40. The above reaction is at equilibrium in a sealed flask. Which of the following actions would cause an increase in the reverse reaction rate after equilibrium has been reestablished?

- (A) Decreasing the temperature
- (B) Adding more $\text{F}_2 (g)$

(C) Removing some $\text{F}_2 (g)$

(D) Removing some $\text{C} (s)$

41. In which of the following molecules does the sulfur atom have the lowest oxidation number?

(A) SO_2

(B) SCl_2

(C) H_2SO_4

(D) SF_6

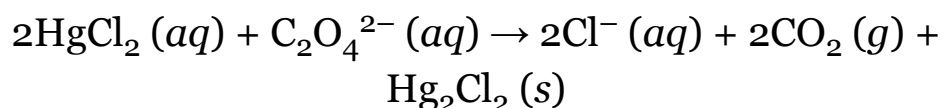
42. At 10°C , the pH of a pure water solution is found to be 7.27. Which of the following best explains why?

(A) Water becomes more basic as temperature decreases.

(B) At 10°C , $[\text{OH}^-] < [\text{H}^+]$.

(C) The auto-ionization of water is endothermic.

(D) The strength of water's intermolecular forces decrease at lower temperatures.



43. Three trials of the above reaction are run with varying concentrations of each reactant and the initial rate of each trial is tracked. Based on the data below, what is the correct rate law for the reaction?

Trial	$[\text{HgCl}_2]$ (M)	$[\text{C}_2\text{O}_4^{2-}]$ (M)	Initial Rate (M/s)
1	0.10	0.10	2.5×10^{-5}

2	0.20	0.10	5.0×10^{-5}
3	0.20	0.20	2.0×10^{-4}

- (A) $\text{Rate} = k[\text{HgCl}_2][\text{C}_2\text{O}_4^{2-}]$
 (B) $\text{Rate} = k[\text{C}_2\text{O}_4^{2-}]^2$
 (C) $\text{Rate} = k[\text{HgCl}_2][\text{C}_2\text{O}_4^{2-}]^2$
 (D) $\text{Rate} = k[\text{HgCl}_2]^2[\text{C}_2\text{O}_4^{2-}]^2$

44. Which of the following ions would have the most unpaired electrons?

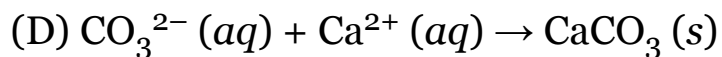
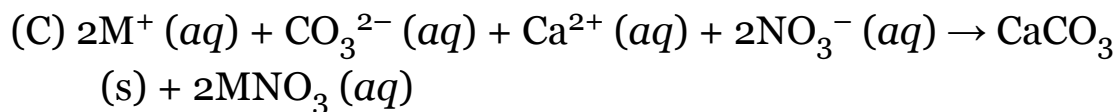
- (A) Na^+
 (B) S^{2-}
 (C) Co^{2+}
 (D) Cr^{2+}

Use the following information to answer questions 45-49.

An 0.740 g sample of an unknown alkali carbonate, identified using M_2CO_3 , is dissolved in 100. mL of water. Excess aqueous $\text{Ca}(\text{NO}_3)_2$ is added to the solution, causing all of the carbonate to precipitate out as calcium carbonate (CaCO_3 , MM = 100 g/mol). The calcium carbonate is filtered, dried, and then massed. It has a final mass of 1.00 g.

45. Identify the correct net ionic equation for the precipitation reaction.

- (A) $\text{M}_2\text{CO}_3 (aq) + \text{Ca}(\text{NO}_3)_2 (aq) \rightarrow \text{CaCO}_3 (s) + 2\text{MNO}_3 (aq)$
 (B) $\text{M}_2\text{CO}_3 (aq) + \text{Ca}(\text{NO}_3)_2 (aq) \rightarrow \text{CaCO}_3 (aq) + 2\text{MNO}_3 (s)$



46. The solution in the beaker after the two solutions are mixed can be best described as

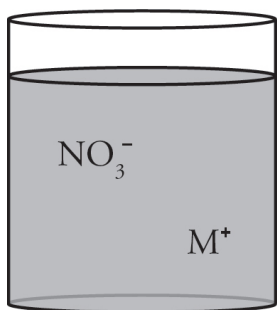
(A) Unsaturated

(B) Saturated

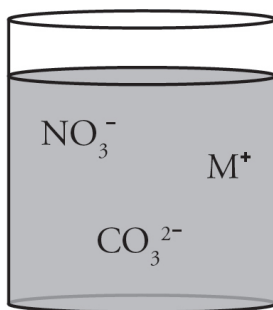
(C) Supersaturated

(D) Hypersaturated

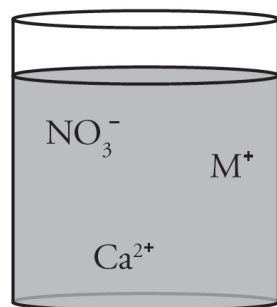
47. Which of the below diagrams correctly shows the ions present in significant amounts after the reaction has gone to completion?



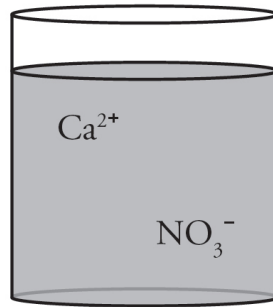
(A)



(C)



(B)



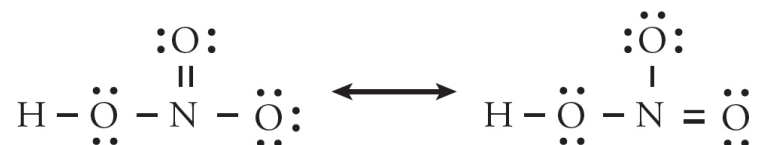
(D)

48. Which of the following is the original alkali carbonate?

- (A) Li_2CO_3
- (B) Na_2CO_3
- (C) K_2CO_3
- (D) Rb_2CO_3

49. Which of the following lab errors would not affect the identification of the alkali cation present in the original sample?

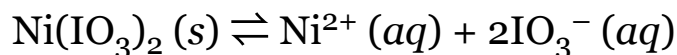
- (A) Spilling some of the alkali carbonate between the balance and the beaker
- (B) Not fully drying the calcium carbonate that is collected at the end
- (C) Leaving a cloudy filtrate after filtering out the precipitate
- (D) Dissolving the alkali carbonate sample in 150 mL of water instead of 100 mL



50. The Lewis diagram for nitric acid, HNO_3 , is above. Which of the following statements is true regarding nitric acid?

- (A) HNO_3 molecules are planar.
- (B) The nitrogen atom carries a negative formal charge.
- (C) All N-O bonds are of identical length.
- (D) There is a third equivalence resonance structure that is not shown.

Use the following information to answer questions 51-53.



Nickel (II) iodate is a slightly soluble salt which dissociates in water via the above equilibrium reaction.

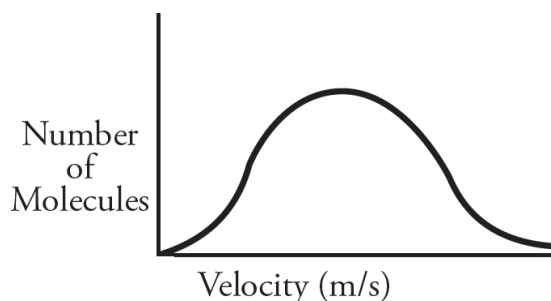
51. In a fully saturated solution of $\text{Ni}(\text{IO}_3)_2$, how do the ion concentrations compare?
- (A) $[\text{Ni}^{2+}] = [\text{IO}_3^-]$
 (B) $[\text{Ni}^{2+}] = 2[\text{IO}_3^-]$
 (C) $2[\text{Ni}^{2+}] = [\text{IO}_3^-]$
 (D) $[\text{Ni}^{2+}]^2 = [\text{IO}_3^-]$
52. In which of the following solutions would the least amount of $\text{Ni}(\text{IO}_3)_2 (s)$ dissolve?
- (A) 0.5 M $\text{Ni}(\text{NO}_3)_2$
 (B) 1.0 M KIO_3
 (C) 2.0 M NaCl
 (D) Pure water
53. If a saturated solution of $\text{Ni}(\text{IO}_3)_2$ is left out overnight and some of the water evaporates, how will that affect the mass of the solid present and/or the concentration of the ions in solution? Assume the temperature of the solution remains constant throughout.

	Mass $\text{Ni}(\text{IO}_3)_2 (s)$	$[\text{Ni}^{2+}]$ and $[\text{IO}_3^-]$
(A)	Increase	Increase
(B)	Increase	Unchanged
(C)	Unchanged	Increase

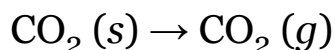
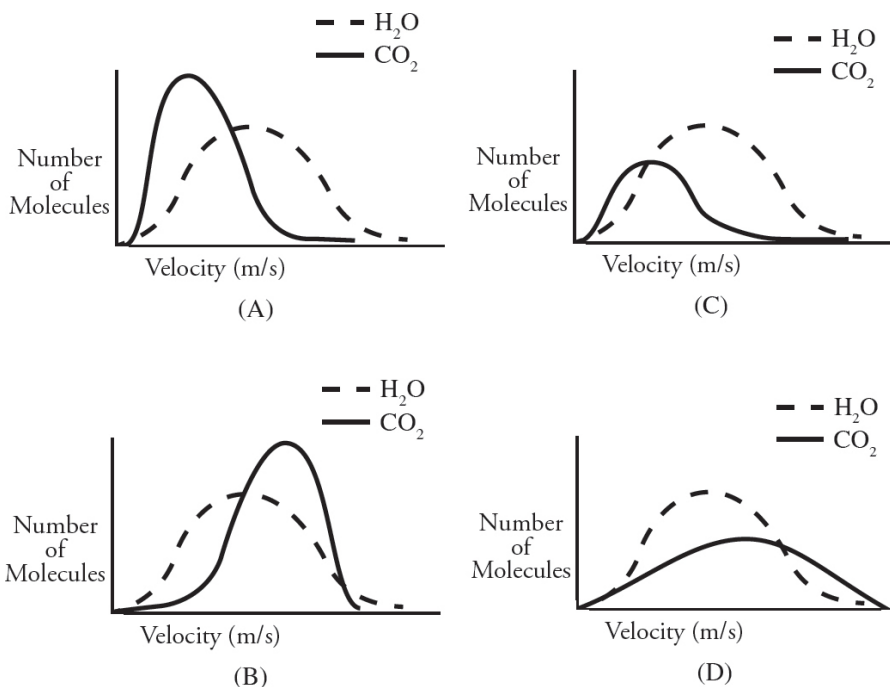
(D)	Unchanged	Unchanged
-----	-----------	-----------

54. A compound containing only hydrogen and carbon is combusted in excess oxygen, producing 4.4 g of CO_2 and 1.8 g of H_2O . Which of the following options could be the molar mass of the compound?

- (A) 18 g/mol
- (B) 28 g/mol
- (C) 38 g/mol
- (D) 48 g/mol

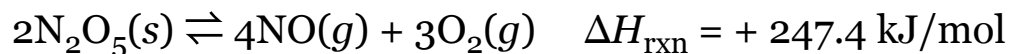


55. The velocity distribution for $\text{H}_2\text{O} (g)$ at a given temperature is drawn above. Which of the following distribution plots would represent the velocity distribution of an equimolar sample of $\text{CO}_2 (g)$ at an identical temperature? The H_2O distribution curve is shown as a dotted line in each option, while the $\text{CO}_2 (g)$ distribution curve is a solid line.



56. Solid carbon dioxide is commonly known by the name dry ice. Dry ice is kept in solid form by sealing it into a high pressure container. When the valve is released, the decrease in pressure causes the CO_2 to sublime via the above reaction. Predict the signs for ΔH° and ΔS° for the sublimation process.

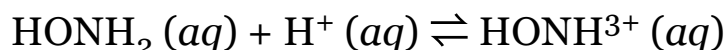
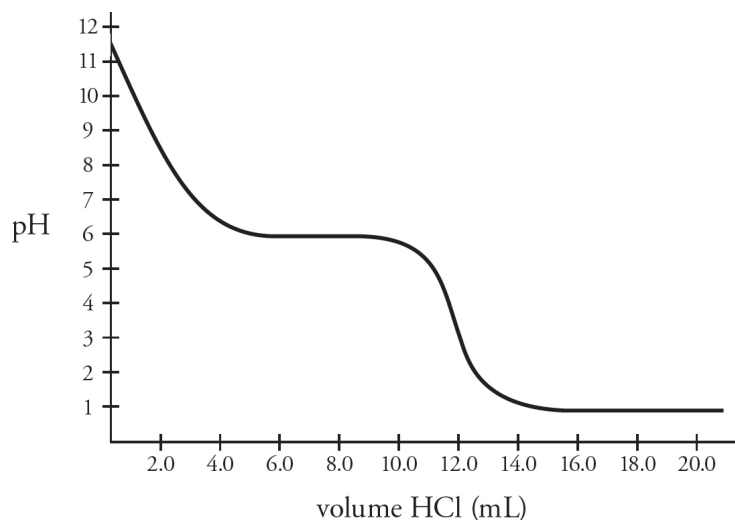
	ΔH°	ΔS°
(A)	+	+
(B)	+	–
(C)	–	+
(D)	–	–



57. Which of the following changes would cause the value for the equilibrium constant for the above reaction to increase?

- (A) Adding some $\text{N}_2\text{O}_5 (s)$
- (B) Increasing the pressure
- (C) Adding some $\text{O}_2 (g)$
- (D) Increasing the temperature

Use the following information to answer questions 58-60.



Hydroxylamine, a weak base, is titrated with 1.0 *M* hydrochloric acid, creating the above titration curve. The temperature of the solution is a constant 25°C.

58. Which of the following reactions best explain why the solution is acidic at the equivalence point of the reaction?
- (A) $\text{HONH}_3^+ (aq) + \text{H}_2\text{O} (l) \rightleftharpoons \text{HONH}_2 (aq) + \text{H}_3\text{O}^+ (aq)$
 - (B) $\text{HONH}_2 (aq) + \text{H}^+ (aq) \rightleftharpoons \text{HONH}_3^+ (aq)$
 - (C) $\text{HONH}_2 (aq) + \text{H}_2\text{O} (l) \rightleftharpoons \text{HONH}_3^+ (aq) + \text{OH}^- (aq)$
 - (D) $\text{HONH}_3^+ (aq) + \text{OH}^- (aq) \rightleftharpoons \text{HONH}_2 (aq) + \text{H}_2\text{O} (l)$

59. What is the approximate pK_b for HONH_2 ?

- (A) 4
- (B) 6
- (C) 8
- (D) 10

60. Which of the following statements correctly identifies the concentration of the various ions in solution after 15.0 mL of HCl has been added?

- (A) $[\text{H}^+] > [\text{HONH}_3^+] > [\text{HONH}_2]$
- (B) $[\text{HONH}_3^+] > [\text{HONH}_2] > [\text{H}^+]$
- (C) $[\text{HONH}_3^+] > [\text{H}^+] > [\text{HONH}_2]$
- (D) $[\text{HONH}_2] > [\text{H}^+] > [\text{HONH}_3^+]$

INFORMATION IN THE TABLE BELOW AND ON THE FOLLOWING PAGES MAY BE USEFUL IN ANSWERING THE QUESTIONS IN THIS SECTION OF THE EXAMINATION.

DO NOT DETACH FROM BOOK.

PERIODIC TABLE OF THE ELEMENTS

PERIODIC TABLE OF THE ELEMENTS																			
1 H 1.008																	2 He 4.00		
3 Li 6.94	4 Be 9.01											5 B 10.81	6 C 12.01	7 N 14.01	8 O 16.00	9 F 19.00	10 Ne 20.18		
11 Na 22.99	12 Mg 24.30											13 Al 26.98	14 Si 28.09	15 P 30.97	16 S 32.06	17 Cl 35.45	18 Ar 39.95		
19 K 39.10	20 Ca 40.08	21 Sc 44.96	22 Ti 47.90	23 V 50.94	24 Cr 52.00	25 Mn 54.94	26 Fe 55.85	27 Co 58.93	28 Ni 58.69	29 Cu 63.55	30 Zn 65.39	31 Ga 69.72	32 Ge 72.59	33 As 74.92	34 Se 78.96	35 Br 79.90	36 Kr 83.80		
37 Rb 85.47	38 Sr 87.62	39 Y 88.91	40 Zr 91.22	41 Nb 92.91	42 Mo 95.94	43 Tc (98)	44 Ru 101.1	45 Rh 102.91	46 Pd 106.42	47 Ag 107.87	48 Cd 112.41	49 In 114.82	50 Sn 118.71	51 Sb 121.75	52 Te 127.60	53 I 126.91	54 Xe 131.29		
55 Cs 132.91	56 Ba 137.33	57 *La 138.91	72 Hf 178.49	73 Ta 180.95	74 W 183.85	75 Re 186.21	76 Os 190.2	77 Ir 192.2	78 Pt 195.08	79 Au 196.97	80 Hg 200.59	81 Tl 204.38	82 Pb 207.2	83 Bi 208.98	84 Po (209)	85 At (210)	86 Rn (222)		
87 Fr (223)	88 Ra 226.02	89 †Ac 227.03	104 Rf (261)	105 Db (262)	106 Sg (266)	107 Bh (264)	108 Hs (277)	109 Mt (268)	110 Ds (271)	111 Rg (272)									
*Lanthanide Series			58 Ce 140.12	59 Pr 140.91	60 Nd 144.24	61 Pm (145)	62 Sm 150.4	63 Eu 151.97	64 Gd 157.25	65 Tb 158.93	66 Dy 162.50	67 Ho 164.93	68 Er 167.26	69 Tm 168.93	70 Yb 173.04	71 Lu 174.97			
			90 Th 232.04	91 Pa 231.04	92 U 238.03	93 Np (237)	94 Pu (244)	95 Am (243)	96 Cm (247)	97 Bk (247)	98 Cf (251)	99 Es (252)	100 Fm (257)	101 Md (258)	102 No (259)	103 Lr (262)			
†Actinide Series																			

ADVANCED PLACEMENT CHEMISTRY EQUATIONS AND CONSTANTS

Throughout the test the following symbols have the definitions specified unless otherwise noted.

L, mL = liter(s), milliliter(s)
 g = gram(s)
 nm = nanometer(s)
 atm = atmosphere(s)

mm Hg = millimeters of mercury
 J, kJ = joule(s), kilojoule(s)
 V = volt(s)
 mol = mole(s)

ATOMIC STRUCTURE

$$E = h\nu$$

$$c = \lambda\nu$$

E = energy
 ν = frequency
 λ = wavelength

Planck's constant, $h = 6.626 \times 10^{-34}$ J s

Speed of light, $c = 2.998 \times 10^8$ m s⁻¹

Avogadro's number = 6.022×10^{23} mol⁻¹

Electron charge, $e = -1.602 \times 10^{-19}$ coulomb

EQUILIBRIUM

$$K_c = \frac{[C]^c[D]^d}{[A]^a[B]^b}, \text{ where } aA + bB \rightleftharpoons cC + dD$$

$$K_p = \frac{(P_C)^c(P_D)^d}{(P_A)^a(P_B)^b}$$

$$K_a = \frac{[H^+][A^-]}{[HA]}$$

$$K_b = \frac{[OH^-][HB^+]}{[B]}$$

$$K_w = [H^+][OH^-] = 1.0 \times 10^{-14} \text{ at } 25^\circ\text{C}$$

$$= K_a \times K_b$$

$$\text{pH} = -\log[H^+], \text{ pOH} = -\log[OH^-]$$

$$14 = \text{pH} + \text{pOH}$$

$$\text{pH} = \text{p}K_a + \log \frac{[A^-]}{[HA]}$$

$$\text{p}K_a = -\log K_a, \text{ p}K_b = -\log K_b$$

Equilibrium Constants

K_c (molar concentrations)

K_p (gas pressures)

K_a (weak acid)

K_b (weak base)

K_w (water)

KINETICS

$$\ln[A]_t - \ln[A]_0 = -kt$$

$$\frac{1}{[A]_t} - \frac{1}{[A]_0} = kt$$

$$t_{1/2} = \frac{0.693}{k}$$

k = rate constant

t = time

$t_{1/2}$ = half-life

GASES, LIQUIDS, AND SOLUTIONS

$$PV = nRT$$

$$P_A = P_{\text{total}} \times X_A, \text{ where } X_A = \frac{\text{moles A}}{\text{total moles}}$$

$$P_{\text{total}} = P_A + P_B + P_C + \dots$$

$$n = \frac{m}{M}$$

$$K = ^\circ\text{C} + 273$$

$$D = \frac{m}{V}$$

$$KE \text{ per molecule} = \frac{1}{2}mv^2$$

Molarity, M = moles of solute per liter of solution

$$A = abc$$

P = pressure

V = volume

T = temperature

n = number of moles

m = mass

M = molar mass

D = density

KE = kinetic energy

v = velocity

A = absorbance

a = molar absorptivity

b = path length

c = concentration

Gas constant, $R = 8.314 \text{ J mol}^{-1} \text{ K}^{-1}$

$= 0.08206 \text{ L atm mol}^{-1} \text{ K}^{-1}$

$= 62.36 \text{ L torr mol}^{-1} \text{ K}^{-1}$

$1 \text{ atm} = 760 \text{ mm Hg}$

$= 760 \text{ torr}$

STP = 273.15 K and 1.0 atm

Ideal gas at STP = 22.4 L mol^{-1}

THERMOCHEMISTRY/ ELECTROCHEMISTRY

$$q = mc\Delta T$$

$$\Delta S^\circ = \sum S^\circ \text{ products} - \sum S^\circ \text{ reactants}$$

$$\Delta H^\circ = \sum \Delta H_f^\circ \text{ products} - \sum \Delta H_f^\circ \text{ reactants}$$

$$\Delta G^\circ = \sum \Delta G_f^\circ \text{ products} - \sum \Delta G_f^\circ \text{ reactants}$$

$$\Delta G^\circ = \Delta H^\circ - T\Delta S^\circ$$

$$= -RT \ln K$$

$$= -nFE^\circ$$

$$I = \frac{q}{t}$$

q = heat

m = mass

c = specific heat capacity

T = temperature

S° = standard entropy

H° = standard enthalpy

G° = standard free energy

n = number of moles

E° = standard reduction potential

I = current (amperes)

q = charge (coulombs)

t = time (seconds)

Faraday's constant, $F = 96,485 \text{ coulombs per mole of electrons}$

$$1 \text{ volt} = \frac{1 \text{ joule}}{1 \text{ coulomb}}$$

CHEMISTRY
Section II

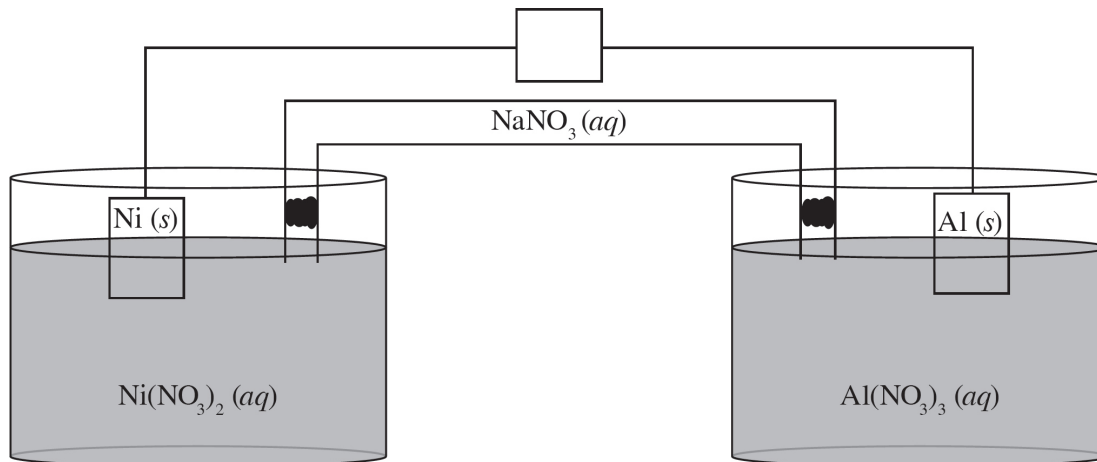
7 Questions

(Total time—105 minutes)

YOU MAY USE YOUR CALCULATOR FOR THIS SECTION

Directions: Questions 1-3 are long free-response questions that require about 23 minutes each to answer and are worth 10 points each. Questions 4-7 are short free-response questions that require about 9 minutes each to answer and are worth 4 points each.

Write your response in the space provided following each question. Examples and equations may be included in your responses where appropriate. For calculations, clearly show the method used and the steps involved in arriving at your answers. You must show your work to receive credit for your answer. Pay attention to significant figures.

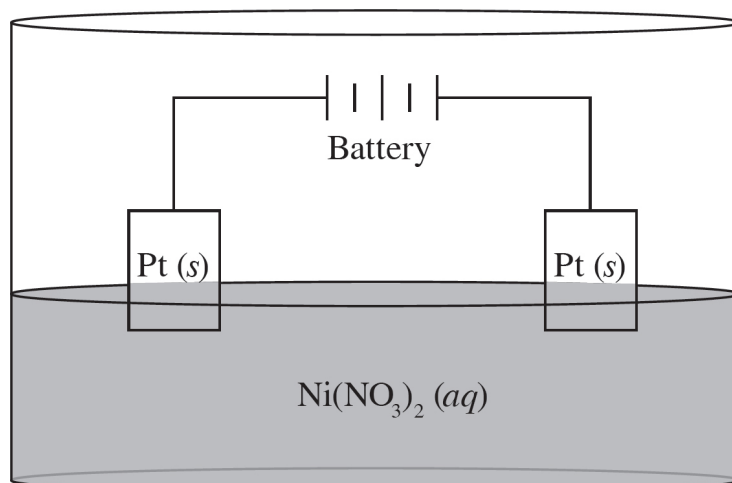


1. A nickel-aluminum voltaic cell is connected as shown above and held at 25°C . The concentration of all solutions is initially at 1.0 M . Some of the standard reduction potentials below may help you in answering the following questions.

Half-Reaction	Standard Reduction Potential (V)
---------------	----------------------------------

$\text{O}_2 (g) + 4\text{H}^+ (aq) + 4e^- \rightarrow 2\text{H}_2\text{O} (l)$	1.23
$\text{Ni}^{2+} (aq) + 2e^- \rightarrow \text{Ni} (s)$	-0.25
$2\text{H}_2\text{O} (l) + 2e^- \rightarrow \text{H}_2 (g) + 2\text{OH}^- (aq)$	-0.83
$\text{Al}^{3+} (aq) + 3e^- \rightarrow \text{Al} (s)$	-1.66

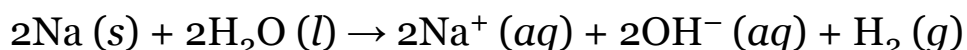
- (a) (i) Write out and balance the full net ionic equation occurring the galvanic cell.
- (ii) Determine E°_{cell} for the reaction in (a)(i).
- (b) If the concentration of the nickel (II) nitrate were to be doubled to 2.0 *M* while the concentration of the aluminum nitrate remained the same, how would that change the E°_{cell} value, if at all? Justify your answer.
- (c) A student studying this cell claims that the electrons flow through the salt bridge from the anode to the cathode. Is this statement correct? Why or why not?



- (d) In a separate experiment, two inert platinum electrodes were placed in a solution of 1.0 *M* nickel (II) nitrate and a

current is run through the electrodes.

- (i) If a current of 2.0 A is run through the nickel (II) nitrate solution for 360. s, how many grams of nickel will plate out?
 - (ii) Will the nickel plate out at the platinum anode, or the platinum cathode? Justify your answer.
 - (iii) Is this reaction thermodynamically favored? Justify your answer.
- (e) When the platinum electrodes are placed in a 1.0 *M* aluminum nitrate and current is run through the solution, no aluminum plates out on either electrode. Why?



2. Like all alkali metals, sodium reacts with water to produce hydrogen gas.

- (a) (i) Which element is being oxidized, and which is being reduced?
 - (ii) What property of alkali metals allows them to react so easily with water?
- (b) (i) If 2.00 g of sodium were to react with 250. mL of pure water at 25°C with an atmospheric pressure of 715 torr, what volume of hydrogen gas would be produced?
- (ii) What would the pH of the remaining solution be after the reaction has gone to completion? Assume the total volume remains unchanged.

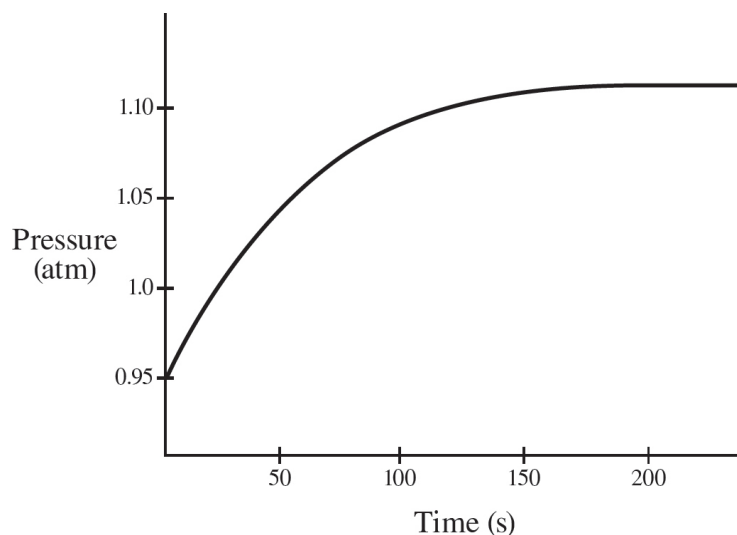


- (c) Three containers are filled completely with identical masses of three different gases, as is shown in the diagram. All containers are held at 25°C .
- (i) In which container, if any, would the pressure be the greatest? Justify your answer.
- (ii) In which container, if any, would the gas molecules have the highest average speed? Justify your answer.
- (d) The three containers are cooled rapidly.
- (i) As the containers are cooled, will the pressure inside them increase, decrease, or stay the same? Justify your answer by discussing the behavior of the gas molecules on a particulate level.
- (ii) Which gas would condense first? Justify your answer.

3.



Potassium chlorate, KClO_3 , will partially decompose at any temperature. 1.00 g of solid potassium chlorate is placed in a 250 mL flask at 23.0°C . The flask is capped, and the pressure inside is monitored until it stabilizes.



- (a) (i) Calculate the partial pressure of the oxygen gas produced by the decomposition of the KClO_3 .
- (ii) Calculate the value for the equilibrium constant expression, K_p , at 23.0°C .
- (b) Would the mass of the KClO_3 increase, decrease, or remain the same between the beginning of the experiment and the point at which equilibrium is established? Justify your answer.
- (c) If the experiment were repeated at the same temperature, but using a 125 mL flask, how would that change the total pressure inside the flask at the end, if at all? Justify your answer.
- (d) Catalysts are often used to lower the activation energy for a given reaction. The experiment is repeated, this time with a MnO_2 catalyst added.
- (i) How do catalysts lower a reaction's activation energy?
- (ii) Using a dashed line, sketch a line on the graph that would indicate any changes that would occur in the data gathered during this trial.

(e) Calculate the value for ΔG for this reaction at 23.0°C .

(f) Is the entropy change that occurs during this reaction positive or negative? Justify your answer.

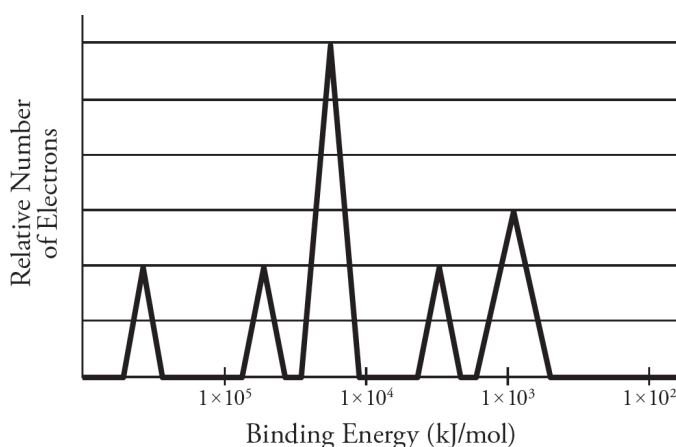
4. The primary acid in lemonade is citric acid ($\text{H}_3\text{C}_6\text{H}_5\text{O}_7$, $\text{p}K_{\text{a}} = 3.08$). To control the pH of many commercially available lemonade drinks, some sodium dihydrogen citrate ($\text{NaH}_2\text{C}_6\text{H}_5\text{O}_7$) is added to create a buffer system.

(a) Write out the reaction which would occur if a strong base were added to a buffered lemonade solution such as the one described above.

(b) What is the mol ratio of $\text{H}_2\text{C}_6\text{H}_5\text{O}_7^- : \text{H}_3\text{C}_6\text{H}_5\text{O}_7$ in a buffered solution with a pH of 3.25?

(c) If a buffered solution of lemonade were to be diluted, would that increase the pH, decrease it, or leave it the same? Justify your answer.

5.



Peak 1	Peak 2	Peak 3	Peak 4	Peak 5
2.7×10^5 kJ/mol	5.4×10^4 kJ/mol	2.1×10^4 kJ/mol	4.9×10^3 kJ/mol	1.0×10^3 kJ/mol

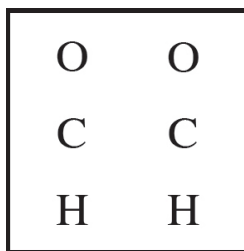
The photoelectron spectrum for a neutral atom is displayed above.

- (a) What is the total number of valence electrons shown on the above spectrum?

A separate sample of the element is bombarded with radiation that has a frequency of $6.3 \times 10^{15} \text{ s}^{-1}$. The energy level is sufficient to only remove the outmost electron.

- (b) How much energy does a single photon of the radiation have?
- (c) How much kinetic energy would the ejected electron have?

6. Glyoxal is an organic compound commonly used in many synthesis reactions.



- (a) In the box below, complete the Lewis electron-dot diagram for the glyoxal molecule by drawing in all electron pairs
- (b) Identify all types of intermolecular forces that are present in a solution of glyoxal.
- (c) Would you expect glyoxal to be miscible with water? Justify your answer.

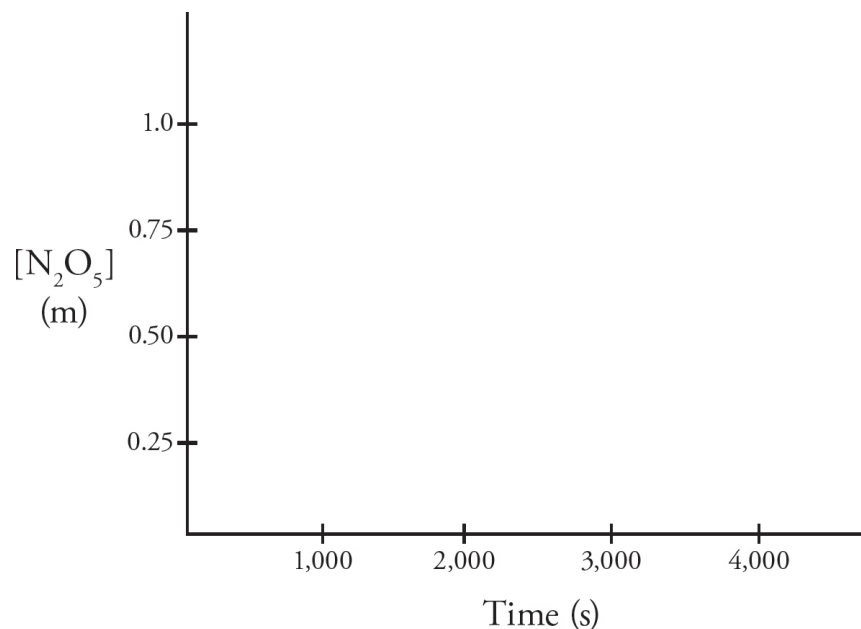
7.



The decomposition of N_2O_5 is a first order reaction that goes to completion. A sample of N_2O_5 gas with a concentration of 1.0 M

is placed in a sealed container at 50°C and allowed to decompose. The half-life of the reaction at 50°C is 1200 seconds.

- (a) On the graph below, draw a graph showing how the concentration of the N_2O_5 sample will change over time.



- (b) Calculate the rate constant for this reaction at 50°C
- (c) Using your graph, determine the instantaneous rate of disappearance for N_2O_5 at $t = 2000$ s.

STOP

END OF EXAM

Practice Test 4: Answers and Explanations

PRACTICE TEST 4: MULTIPLE-CHOICE ANSWER KEY

- | | | |
|-------|-------|-------|
| 1. C | 21. B | 41. B |
| 2. A | 22. B | 42. C |
| 3. B | 23. A | 43. C |
| 4. B | 24. B | 44. D |
| 5. D | 25. C | 45. D |
| 6. B | 26. D | 46. B |
| 7. D | 27. D | 47. B |
| 8. C | 28. A | 48. A |
| 9. D | 29. D | 49. D |
| 10. A | 30. A | 50. A |
| 11. A | 31. C | 51. C |
| 12. B | 32. A | 52. B |
| 13. C | 33. D | 53. B |
| 14. A | 34. C | 54. B |
| 15. A | 35. A | 55. D |
| 16. C | 36. A | 56. A |
| 17. A | 37. B | 57. D |
| 18. D | 38. A | 58. A |
| 19. D | 39. D | 59. C |
| 20. B | 40. B | 60. C |

SECTION I—MULTIPLE-CHOICE ANSWERS AND EXPLANATIONS

1. C

When the Ag_2CO_3 dissociates, it produces two Ag^+ ions for every one CO_3^{2-} ion.

2. A

Henderson-Haselbalch is used here, so $\text{pH} = \text{p}K_a + \log [\text{NO}_2^-]/[\text{HNO}_2]$. For the pH to be the same as the $\text{p}K_a$ of the acid, the logarithm term must cancel out. That happens when $[\text{NO}_2^-] = [\text{HNO}_2]$.

3. B

The graph indicates the decomposition reaction has a constant half-life. This means the reaction must be first order.

4. B

There is resonance between the two S-O bonds where the H is not attached to the other side of the oxygen atom. These bonds are identical in length, but would be shorter than the true single S-O bond with the oxygen that does have the hydrogen bonded to it.

5. D

The H^+ ions are being reduced, but the K (s) is being oxidized, so the potassium half-reaction has to be flipped.

It also has to be multiplied by two so the electrons will cancel out when the two half reactions are combined.

6. **B**

Vapor pressure is a function of the strength of the intermolecular forces. When the methanol is heated, the methanol molecules will be moving faster, meaning they are more likely to have enough energy to break free of the IMFs, meaning more of them will end up in a gaseous state, increasing the vapor pressure.

7. **D**

First solve for the heat lost by the water using $q = mc\Delta T$. In this case, the mass is the mass of the dissolved compound plus that of the water, so about 100 g.

$$q = (100 \text{ g})(4.2 \text{ J/g}^\circ\text{C})(-3.0^\circ\text{C}) \quad q = -1260 \text{ J}$$

To determine the enthalpy of reaction, flip the sign and then divide by the number of moles.

$$\Delta H = q_{\text{rxn}}/n \quad \Delta H = 1260 \text{ J}/0.10 \text{ mol} = 12600 \text{ J/mol} = 12.6 \text{ kJ/mol}$$

8. **C**

The reaction occurs, so it must be favored. The surrounding water lost energy, meaning the reaction required energy to occur. This means the reaction is endothermic, which is not a favorable enthalpy change. The KNO_2 starts as a solid and ends up as dissociated ions. Those ions are much more disordered than the solid, meaning disorder increases, which is a favorable entropy change.

9. **D**

When an ionic salt dissolves in water, the energy required is that to break up the ionic lattice, whereas the energy released comes from the attraction of the ions to the dipoles of the surrounding water molecules. The reaction was endothermic, which indicates the lattice energy is greater than the ion-dipole attractions. (Note that the reaction still proceeds due to its favorable entropy change!)

10. **A**

If not all of the salt dissolved, the moles of salt in the denominator of the enthalpy of reaction expression would be artificially high, making the enthalpy of reaction artificially low.

11. **A**

For the salt to precipitate, the value for the reaction quotient has to be equal to or greater than the equilibrium constant for the reaction.

$$Q_{\text{sp}} = [\text{Sr}^{2+}][\text{IO}_3^-]^2 \quad 1.0 \times 10^{-7} = [\text{Sr}^{2+}](1.0 \times 10^{-2})^2 \quad [\text{Sr}^{2+}] = 1.0 \times 10^{-3} \text{ M}$$

12. **B**

After all of an acid has reacted during a titration, all that is present in the solution is the conjugate base of that acid. In a polyprotic titration, the endpoint occurs after all of the protons have been neutralized. In each case, the lower the K_a is for the final dissociation of the weak acid, the higher the K_b will be of its conjugate base (because for any conjugate pair, $K_a \times K_b = 1.0 \times 10^{-14}$). A higher K_b means a stronger conjugate base, leading to a higher pH at the endpoint.

13. **C**

A catalyst speeds up a reaction by lowering the activation energy of that reaction. The activation energy is shown on the graph as the height of the “hump,” so a catalyzed reaction will have a lower hump. Note that the enthalpy change of the reaction (the distance between the energy level of the reactants and the products) will remain unchanged.

14. **A**

The higher the K_a value, the greater the percent dissociation for the acid is, and the higher the concentration of H^+ ions will be.

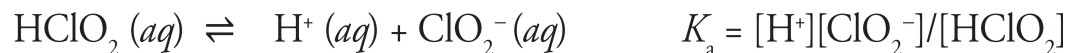
15. **A**

To determine the conjugate base of a weak acid, simply remove the proton (H^+).

16. **C**

To create a buffer, similar amounts of a weak acid and its conjugate must be present. The conjugate base of $HClO_2$ is ClO_2^- , so adding a soluble salt with the ClO_2^- ion present would create a buffer.

17. **A**



$$1.0 \times 10^{-2} = x^2/1.0 - x \quad x = 1.0 \times 10^{-2} M = [H^+]$$

$$-\log[H^+] = pH \quad -\log(1.0 \times 10^{-2}) = 2$$

Note that the above calculations ignore the “ $-x$ ” term in the denominator. While HClO_2 has a high enough K_a that the dissociation will be significant and affect the pH, it can be still be ignored for the purposes of approximation.

18. **D**

The first dissociation is $\text{H}_2\text{CO}_3 \rightleftharpoons \text{H}^+ + \text{HCO}_3^-$. The second is $\text{HCO}_3^- \rightleftharpoons \text{H}^+ + \text{CO}_3^{2-}$. Note that H^+ is a product of both reactions. After the first H^+ dissociates off of the H_2CO_3 molecule, that will cause a left shift in the second dissociation reaction, meaning less HCO_3^- will be able to dissociate.

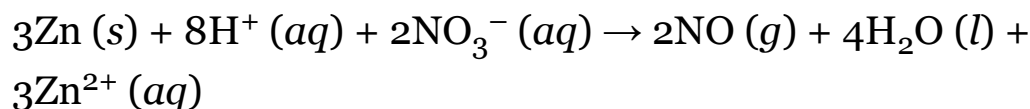
19. **D**

Formal charge is calculated by subtracting the number of assigned electrons on an atom in a Lewis diagram from the number of its valence electrons. When looking at an atom, lone pairs are considered to be two assigned electrons, and any type of bond is considered to be one (so a single bond is one, a double bond would be two, etc). Constructing a formal charge chart for the identified atoms shows that of the identified atoms only O_z would have a negative formal charge.

	P	O_x	O_y	O_z
Valence	5	6	6	6
Assigned	5	6	6	7
Formal Charge	0	0	0	-1

20. **B**

To fully balance the reaction, the charge must be balanced. This required multiplying the zinc oxidation half-reaction by two, and the nitrate reduction half-reaction by three. Doing that and combining the reactions yields:



21. **B**

Orbital hybridization can be determined by looking at the number of charge clouds around the atom in question. The oxygen has three charge clouds—two lone pair, and one bond. That is consistent with sp^2 hybridization.

22. **B**

Single bonds are longer than double bonds, and when comparing the S-C and C-H bonds, a sulfur atom is bigger than a hydrogen atom, which would mean the S-C bond is longer.

23. **A**

Oxygen is very electronegative, and would be the most effective at attracting electrons toward it. That would give it a negative dipole, and since oxygen is directly bonded to sulfur, sulfur would have a positive dipole.

24. **B**

In order to separate out the propanol, it must be boiled, so the temperature must be above the boiling point of propanol. However, the temperature should also be below the boiling point of propionic acid, because if both substances boil they will not separate effectively. There are two options in the temperature range, and the lower one is

better because the propionic acid will have a lower vapor pressure at 105°C, meaning less of it will evaporate and contaminate the distillate.

25. **C**

For any reaction, the coefficients on the reactants in the slowest elementary step are equal to the order of those reactants in the rate law. So, the rate law for the given mechanism is $\text{rate} = k[\text{NO}_2]^2$. A second-order reactant produces a straight line graph when graphing the reciprocal of that reactant vs. time.

26. **D**

The type of radiation that is used to study sub-atomic particles depends on the size of the particles. The higher the frequency of the radiation, the smaller the particles that can be studied. For studying electrons, ultraviolet radiation is often used. Studying bonds (shared pairs of electrons) generally uses infrared radiation, while studying entire molecules uses microwave radiation.

27. **D**

The diagrams can be diagnosed the relative atomic sizes of the ions. Li^+ and F^- would be the smallest ions, which are present in diagram Z. Na^+ and Cl^- would be the next biggest, so diagram X. Finally, K^+ and Br^- would be the largest, which corresponds with diagram Y.

28. **A**

All three compounds contain ions of identical charge, and the only other factor that can affect the lattice energy (a type of Coulombic energy) is size, which is inversely

proportional. Li^+ and F^- are the smallest ions, meaning they have the greatest amount of lattice energy and the highest melting point.

29. **D**

When the cations lose their only valence electron, the valence shell becomes the one beneath it. For instance, Na^+ would have a configuration of $1s^2 2s^2 2p^6$. When the halide anions gain an electron, they too complete their outmost p -subshell. Li^+ is the exception because after lithium loses an electron, all that is left is the first energy level, which has no p -subshell.

30. **A**

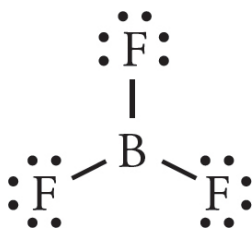
During a liquid to gas phase change, any energy added goes into breaking the intermolecular forces between molecules. The temperature of a boiling solution remains constant, and thus, so does the speed of the molecules.

31. **C**

A fluoride atom is $[\text{He}]2s^2 2p^5$. A fluorine ion would be $[\text{He}]2s^2 2p^6$. The $2p$ peak is the rightmost peak, and in the ion it would contain one more electron and thus be higher than the $2p$ peak from the neutral atom.

32. **A**

Boron has a very low electronegativity value for a non-metal, and within a molecule it is considered stable with just six valence electrons. Thus, a molecule of BF_3 would be stable with three single bonds and would look like this:



33. **D**

There are nine oxygen atoms present in the hydrate: four in the anhydrous CuSO_4 salt, and five more from the five water molecules. In one mole of the hydrate, the mass of those oxygen atoms would be $16(9) = 144$ g. The oxygen thus represents 144 g out of a 250 g sample. It would maintain the same mass ratio in a 25.0 g sample, which would be 14.4 g.

34. **C**

At STP, 1 mol of any gas has a volume of 22.4 L. If the gas in questions has a volume of 5.6 L, that represents $5.6/22.4 = 0.25$ moles of the gas. Multiplying the mass of the sample by four would then yield the mass of 1.0 mole of the gas. $10.0 \text{ g} \times 4 = 40.0 \text{ g/mol}$, which is the molar mass of argon.

35. **A**

The coefficients on the reactants present in the slowest elementary step become the order of those reactants in the overall reaction. Therefore, the first step must be the slow/rate-determining step. Molecularity is dependent on how many reactant molecules there are, which in this case is just one.

36. **A**

All of the potential options use seconds, so the rate must be measured in M/s. With that in mind, it's just a matter of

using dimensional analysis on the rate law. Rate =
 $k[\text{NO}_2\text{Cl}] \text{ M/s} = k(\text{M}) \text{ k} = \text{s}^{-1}$

37. **B**

Catalysts are present in the reactants of the first step and the products of the final step. No species in this reactant fits that profile. Intermediates are present in the products of an early step and the reactants of a later step. Cl fits the bill.

38. **A**

The overall reaction is exothermic, meaning the final energy level of the products must be lower than the initial energy level of the reactants. Additionally, step 1 of the process must be endothermic, since bonds are broken in step 1 without any new bonds being formed. The first hump is for the first step, so after that hump the energy level must be higher than where it started.

39. **D**

$$2.80 \text{ g CO} \times \frac{1 \text{ mol CO}}{28.0 \text{ g CO}} \times \frac{1 \text{ mol}_{\text{rxn}}}{3 \text{ mol CO}} \times \frac{26.4 \text{ kJ}}{1 \text{ mol}_{\text{rxn}}} = 0.88 \text{ kJ}.$$

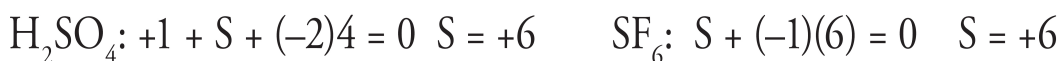
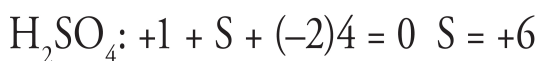
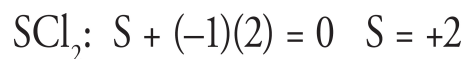
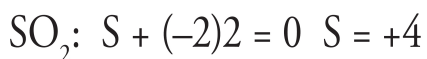
The positive sign means the reaction is endothermic, and thus heat was absorbed.

40. **B**

Adding the $\text{F}_2(g)$ would cause an immediate shift to the right, which would increase the forward reaction rate. However, the key is that after equilibrium is reestablished there is more matter in the flask, meaning both the forward

and reverse reaction rates will be higher than they were initially (and still equal to each other).

41. **B**



42. **C**

Pure water has a neutral pH of 7 at 25°C only. When the temperature changes, so does the pH of water. The auto-ionization of water is $\text{H}_2\text{O} (l) \rightleftharpoons \text{H}^+ (aq) + \text{OH}^- (aq)$. When temperature decreases, that equilibrium shifts to the left, as it would in any endothermic process. This decreases the value for the equilibrium constant, meaning that both $[\text{H}^+]$ and $[\text{OH}^-]$ decrease (but still remain the same). If $[\text{H}^+]$ and $[\text{OH}^-]$ decrease, the pH of the water increases, as $[\text{H}^+]$ and pH are inversely proportional.

43. **C**

Between trials 1 and 2, the $[\text{HgCl}_2]$ doubles while $[\text{C}_2\text{O}_4^{2-}]$ stays the constant. The rate of the reaction also doubles, which demonstrates a first order relationship with regards to $[\text{HgCl}_2]$. Between trials 2 and 3, $[\text{C}_2\text{O}_4^{2-}]$ doubles while $[\text{HgCl}_2]$ remains the same. The rate increases by a factor of four, demonstrating a second order relationship with regards to $[\text{C}_2\text{O}_4^{2-}]$.

44. **D**

Choices (A) and (B) have no unpaired electrons. Comparing (C) and (D), both are transition metals, which always lose their s electrons first when forming ions. Cr^{2+} would thus

have four unpaired *d* electrons, while Co^{2+} would only have three.

45. **D**

In a net ionic reaction, the only ions that appear are those that undergo a chemical change. In this case, that is the Ca^{2+} and CO_3^{2-} ions. The M^+ and NO_3^- are spectator ions and would not appear in the net ionic equation.

46. **B**

Any solution that has a solid precipitate at the bottom is considered to be saturated at the very least. For it to be supersaturated, the solution would need to be heated to dissolve more precipitate and then cooled back to its original temperature.

47. **B**

The calcium nitrate is described as being in excess, so there will definitely be Ca^{2+} and NO_3^- ions in solution. In addition, the M^+ alkali cation does not react, so it will also be present in the final solution.

48. **A**

$$1.00 \text{ g CaCO}_3 \times \frac{1 \text{ mol CaCO}_3}{100. \text{ g CaCO}_3} = 0.010 \text{ mol}$$

$$\text{CaCO}_3 \times \frac{1 \text{ mol CO}_3^{2-}}{1 \text{ mol CaCO}_3} = 0.010 \text{ mol CO}_3^{2-}$$

$$0.010 \text{ mol CO}_3^{2-} \times \frac{1 \text{ mol M}_2\text{CO}_3}{1 \text{ mol CO}_3^{2-}} = 0.010 \text{ mol M}_2\text{CO}_3$$
$$\frac{0.74 \text{ g}}{0.010 \text{ mol}} = 74 \text{ g/mol}$$

74 g/mol is the molar mass of lithium carbonate.

49. **D**

If the alkali carbonate dissolves fully in 100. mL of water, it will also dissolve fully in 150. mL of water. The same amount of carbonate will be in solution, leading to no difference in the final mass of the CaCO_3 precipitate.

50. **A**

The resonance shown here is for the N-O bonds that do not have hydrogen attached on the other end of the oxygen. This means that two of the N-O bonds are identical (bond order = 1.5), but the third one (from N-O-H) is a true single bond. The geometry around the N is trigonal planar, and the geometry around the O in the N-O-H is bent, both of which are planar geometries.

51. **C**

Based on the balanced equilibrium reaction, in a saturated solution there will be twice as many IO_3^- ions as there are Ni^{2+} ions.

52. **B**

The common ion effects states that if the salt that is being dissolved shares an ion with the solution that it is being dissolved in, less of that salt will dissolve. This is because the presence of that ion will push the solubility equilibrium to the left. Both (A) and (B) share ions with $\text{Ni}(\text{IO}_3)_2$, but there are more of those ions present in (B) due to the higher concentration.

53. **B**

As long as temperature stays unchanged, so does the K_{sp} value for the salt, meaning the ion concentration will not change. However, if some water evaporates, that means the volume decreased. For the ion concentration to remain unchanged, a decrease in volume must be accompanied by a decrease in mass of the ions dissolved in solution. Those ions will recombine and precipitate out, increasing the mass of solid present at the bottom.

54. **B**

For any hydrocarbon combustion, all of the carbon in the original compound ends up in CO_2 , and all of the hydrogen ends up in H_2O . Some stoichiometry gives us the mole ratio of carbon to hydrogen in the original compound.

$$4.4 \text{ g CO}_2 \times \frac{1 \text{ mol CO}_2}{44.0 \text{ g CO}_2} \times \frac{1 \text{ mol C}}{1 \text{ mol CO}_2} = 0.10 \text{ mol C}$$

$$1.8 \text{ g H}_2\text{O} \times \frac{1 \text{ mol H}_2\text{O}}{18.0 \text{ g H}_2\text{O}} \times \frac{2 \text{ mol H}}{1 \text{ mol H}_2\text{O}} = 0.20 \text{ mol H}$$

Thus, the ratio of C:H in the compound must be 1:3. A single CH_3 unit would have a mass of 15.0 g/mol, and two of those units (so, C_2H_6) would have a molar mass of 30 g/mol.

55. **D**

CO_2 has a greater molar mass than H_2O . At identical temperatures, both gases have identical kinetic energy. Given that $\text{KE} = \frac{1}{2}mv^2$, if the CO_2 has a higher mass it must have a lower average velocity, which is why the apex is lower. Additionally, the area under both of the curves

represents the number of molecules and must be the same given that the samples are equimolar, which is why the (lower) apex must shift right.

56. **A**

The solid turning into a gas is an increase in disorder, making ΔS° positive. In order for the solid to turn into a gas, IMFs must be broken, which requires the input of energy from the environment. So, ΔH° is also positive.

57. **D**

The only way to change the equilibrium constant for any reaction is by changing the temperature. Increasing the temperature on an endothermic reaction permanently shifts it right, creating more products and increasing the value of the equilibrium constant.

58. **A**

At the equivalence point, all of the HONH_2 has been protonated by the H^+ because equal amounts have been added. The only ion present at this point is the HONH_3^+ , which reacts with water to create an acidic environment.

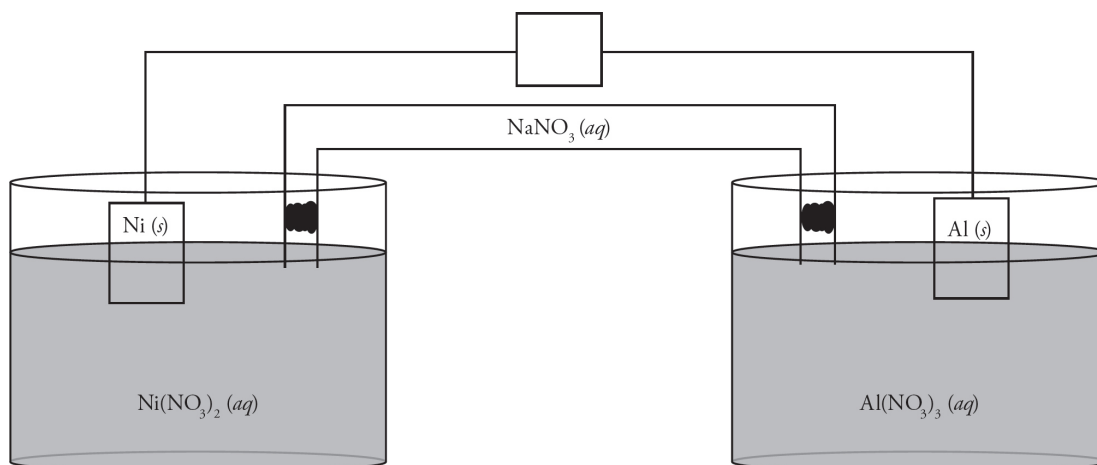
59. **C**

At the half-equivalence point of this reaction, the pOH of the solution is equal to the $\text{p}K_b$ of hydroxylamine. This occurs at about 6.0 mL, at which point the pH is 6. That means the pOH is 8 (because at 25°C $\text{pH} + \text{pOH} = 14$), as is the $\text{p}K_b$ for hydroxylamine.

60. **C**

15.0 mL is past the equivalence point, so all of the original HONH_2 has been protonated and there is none left in solution. It took 12.0 mL of H^+ to create the HONH_3^+ in solution, and after that, only 3.0 mL of H^+ have been added. So, the amount of HONH_3^+ is still greater than the amount of unreacted H^+ in solution.

SECTION II—FREE-RESPONSE ANSWERS AND EXPLANATIONS

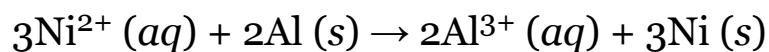


1. A nickel-aluminum voltaic cell is connected as shown above and held at 25°C. The concentration of all solutions is initially at 1.0 *M*. Some of the standard reduction potentials below may help you in answering the following questions.

Half-Reaction	Standard Reduction Potential (V)
$\text{O}_2(g) + 4\text{H}^+(aq) + 4e^- \rightarrow 2\text{H}_2\text{O}(l)$	1.23
$\text{Ni}^{2+}(aq) + 2e^- \rightarrow \text{Ni}(s)$	−0.25
$2\text{H}_2\text{O}(l) + 2e^- \rightarrow \text{H}_2(g) + 2\text{OH}^-(aq)$	−0.83
$\text{Al}^{3+}(aq) + 3e^- \rightarrow \text{Al}(s)$	−1.66

- (a) (i) Write out and balance the full net ionic equation occurring the galvanic cell.

The trick here is to make sure the electrons are balanced in the two half-reactions. This requires multiplying the nickel half-reaction by 3, and the aluminum half-reaction by 2. When you combine them, 6 moles of electrons will then cancel.



- (ii) Determine E°_{cell} for the reaction in (a)(i).

$$-0.25 \text{ V} - (-1.66 \text{ V}) = 1.41 \text{ V}$$

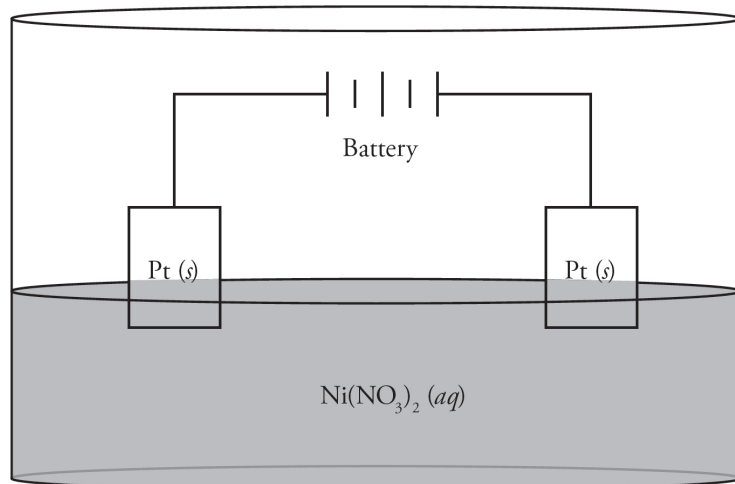
- (b) If the concentration of the nickel (II) nitrate were to be doubled to 2.0 *M* while the concentration of the aluminum nitrate remained the same, how would that change the E°_{cell} value, if at all? Justify your answer.

Whether or not the voltage increases or decreases depends on if the change described brings the reaction closer to or further from equilibrium. To determine that, look at *Q*.

$$Q = \frac{[\text{Al}^{3+}]^2}{[\text{Ni}^{2+}]^3}$$

If the nickel concentration increases while the aluminum concentration remains the same, that means *Q* will decrease. As this cell is voltaic, that means *K* is much, much bigger than one. If *Q* is getting smaller, that it taking the cell further from equilibrium. For any cell, at equilibrium the voltage is zero, so taking the cell further from equilibrium takes the voltage further from zero. Thus, E°_{cell} would increase.

- (c) A student studying this cell claims that the electrons flow through the salt bridge from the anode to the cathode. Is this statement correct? Why or why not?



This statement is incorrect. Electrons do not flow through the salt bridge. Instead, they flow through the wires.

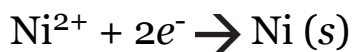
- (d) In a separate experiment, two inert platinum electrodes were placed in a solution of 1.0 *M* nickel (II) nitrate and a current is run through the electrodes.
- (i) If a current of 2.0 A is run through the nickel (II) nitrate solution for 360. s, how many grams of nickel will plate out?

Electroplating problems are all about dimensional analysis/unit conversions. Remember, an Ampere is equivalent to a Coulomb per second.

$$360. \text{ s} \times \frac{2.0 \text{ C}}{\text{s}} \times \frac{1 \text{ mol } e^-}{96500 \text{ C}} \times \frac{1 \text{ mol Ni}}{2 \text{ mol } e^-} \times \frac{58.69 \text{ g}}{1 \text{ mol Ni}} = 0.22 \text{ g Ni}$$

- (ii) Will the nickel plate out at the platinum anode, or the platinum cathode? Justify your answer.

It will plate out at the cathode, because the cathode is where the reduction reaction occurs.

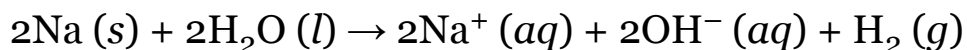


- (iii) Is this reaction thermodynamically favored? Justify your answer.

This reaction is not favored, as an outside source of energy (in this case, electrical current) is required for the reaction to occur.

- (e) When the platinum electrodes are placed in a 1.0 *M* aluminum nitrate and current is run through the solution, no aluminum plates out on either electrode. Why?

The reduction potential of the Al^{3+} ion, -1.66 V , is lower than that of water, -0.83 V . Thus, when current is run through the solution, the reduction of water will occur. This will produce hydrogen gas as water is reduced, but the aluminum ions will stay in solution.



2. Like all alkali metals, sodium reacts with water to produce hydrogen gas.

- (a) (i) Which element is being oxidized, and which is being reduced?

Sodium is being oxidized, going from an oxidation state of 0 to +1. Hydrogen is being reduced, going from a state of +1 in water to 0 in hydrogen gas.

- (ii) What property of alkali metals allows them to react so easily with water?

Alkali metals all have very low ionization energies, and so they can be readily oxidized by water.

- (b) (i) If 2.00 g of sodium were to dissolve in 250. mL of pure water at 25°C with an atmospheric pressure of 715 torr, what volume of hydrogen gas would be produced?

First, figure out how many moles of hydrogen gas are being produced.

$$2.00 \text{ g Na} \times \frac{1 \text{ mol Na}}{22.99 \text{ g}} \times \frac{1 \text{ mol H}_2}{2 \text{ mol Na}} = 0.0435 \text{ mol H}_2$$

Then, use the ideal gas law to convert that to volume.

$$P = \frac{715 \text{ torr}}{760 \text{ torr}} = 0.941 \text{ atm} \quad T = 25^\circ\text{C} + 273 = 298 \text{ K}$$

$$PV = nRT$$

$$PV = nRT$$

$$(0.941)(V) = (0.0435)(0.0821)(298)$$

$$V = 1.13 \text{ L}$$

- (ii) What would the pH of the remaining solution be after the reaction has gone to completion? Assume the total volume remains unchanged.

The pH will be dependent on the concentration of the OH⁻ ions, as they are a strong base.

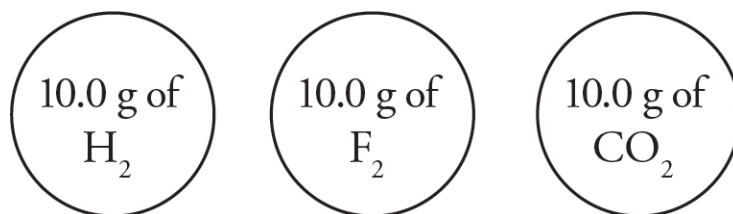
$$2.00 \text{ g Na} \times \frac{1 \text{ mol Na}}{22.99 \text{ g}} \times \frac{2 \text{ mol OH}^-}{2 \text{ mol Na}} = 0.0870 \text{ mol OH}^-$$

$$\frac{0.0870 \text{ mol OH}^-}{0.250 \text{ L}} = 0.348 \text{ M}$$

$$-\log[\text{OH}^-] = \text{pOH} \quad -\log(0.348) = 0.458$$

$$\text{pH} + \text{pOH} = 14 \quad \text{pH} + 0.458 = 14$$

$$\text{pH} = 13.54$$



(c) Three containers are filled completely with identical masses of three different gases, as is indicated on the diagram. All containers are held at 25°C.

(i) In which container, if any, would the pressure be the greatest? Justify your answer.

If the same mass of gas is present in each container, the gas with the lowest molar mass would have the greatest number of moles present. Moles and pressure are directly proportional, so in this case the container with the hydrogen gas would have the greatest pressure.

(ii) In which container, if any, would the gas molecules have the highest average speed? Justify your answer.

As all containers are at the same temperature, the gases inside have the same amount of kinetic energy. Both the mass and velocity of the individual gas molecules contribute to their total kinetic energy. If the kinetic energy is the same, the gas molecules with the lowest mass would also have the highest velocity to compensate.

Thus, the hydrogen molecules would have the greatest average velocity.

(d) The three containers are cooled rapidly.

(i) As the containers are cooled, will the pressure inside them increase, decrease, or stay the same? Justify your answer by discussing the behavior of the gas molecules on a particulate level.

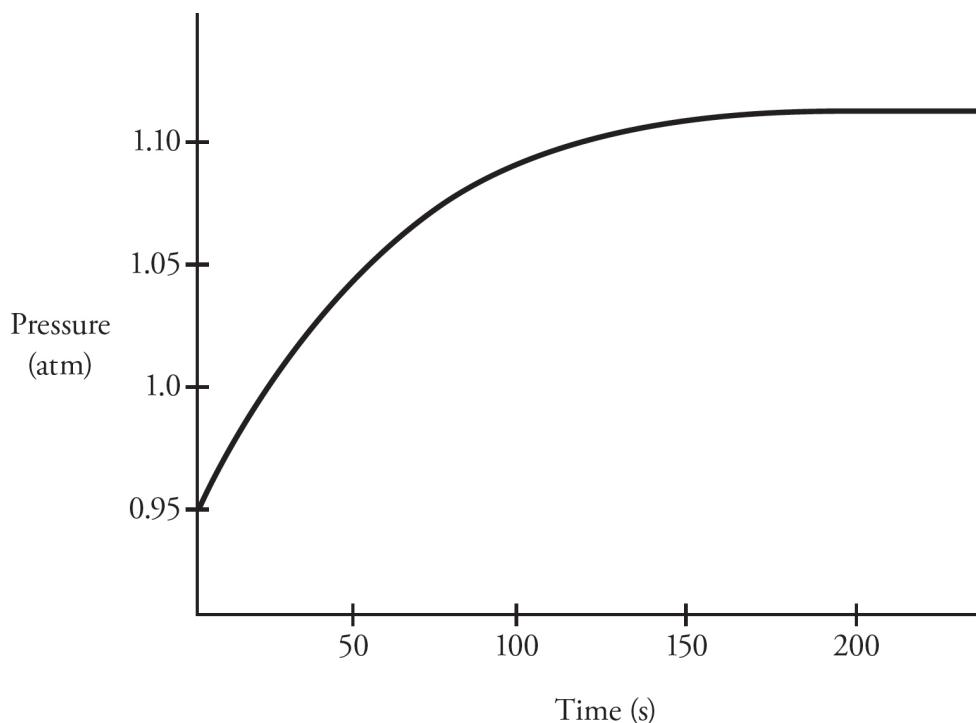
The pressure in each container will decrease. As the temperature drops, the molecules in each container will slow down. They will thus hit the sides of the container less often and with less force, meaning the total pressure decreases.

(ii) Which gas would condense first? Justify your answer.

A phase change from gas to liquid will occur once the gas molecules are moving slow enough that the intermolecular forces between them become significant enough to affect their behavior. The gas with the strongest IMFs will condense first. As all three gases are non-polar, the only IMFs present are London dispersion forces. The CO_2 gas has the most polarizable electron cloud, and so it has the strongest LDFs and would condense first.



3. Potassium chlorate, KClO_3 , will partially decompose at any temperature. 1.00 g of solid potassium chlorate is placed in a 250 mL flask at 23.0°C . The flask is capped, and the pressure inside is monitored until it stabilizes.



- (a) (i) Calculate the partial pressure of the oxygen gas produced by the decomposition of the KClO_3 .

The initial pressure of the flask at $t = 0$ s is the pressure of just the atmosphere. The total pressure at the end consists of both the atmosphere and the produced oxygen gas. Subtracting the two values would give you the partial pressure of the oxygen gas.

$$1.11 \text{ atm} - 0.95 \text{ atm} = 0.16 \text{ atm} = P_{\text{O}_2}$$

- (ii) Calculate the value for the equilibrium constant expression, K_p , at 23.0°C .

Remember, solids do not appear in equilibrium expressions.

$$K_p = (P_{\text{O}_2})^3 =$$

$$K_p = (0.16)^3 = 4.1 \times 10^{-3}$$

- (b) Would the mass of the KClO_3 increase, decrease, or remain the same between the beginning of the experiment and the point at which equilibrium is established? Justify your answer.

The mass of the KClO_3 would decrease. The reason that solids do not appear in the equilibrium expression is because their concentration is unchanging. However, the mass of the KClO_3 can still decrease without changing the concentration. For any products to be produced, at least some of the KClO_3 must decompose, reducing its mass.

- (c) If the experiment were repeated at the same temperature, but using a 125 mL flask, how would that change the total pressure inside the flask at the end, if at all? Justify your answer.

The pressure in the flask would be the same. The initial atmospheric pressure would be the same, and as long as temperature is constant the value for K_p would not change. If the K_p value does not change, neither does the partial pressure of the oxygen.

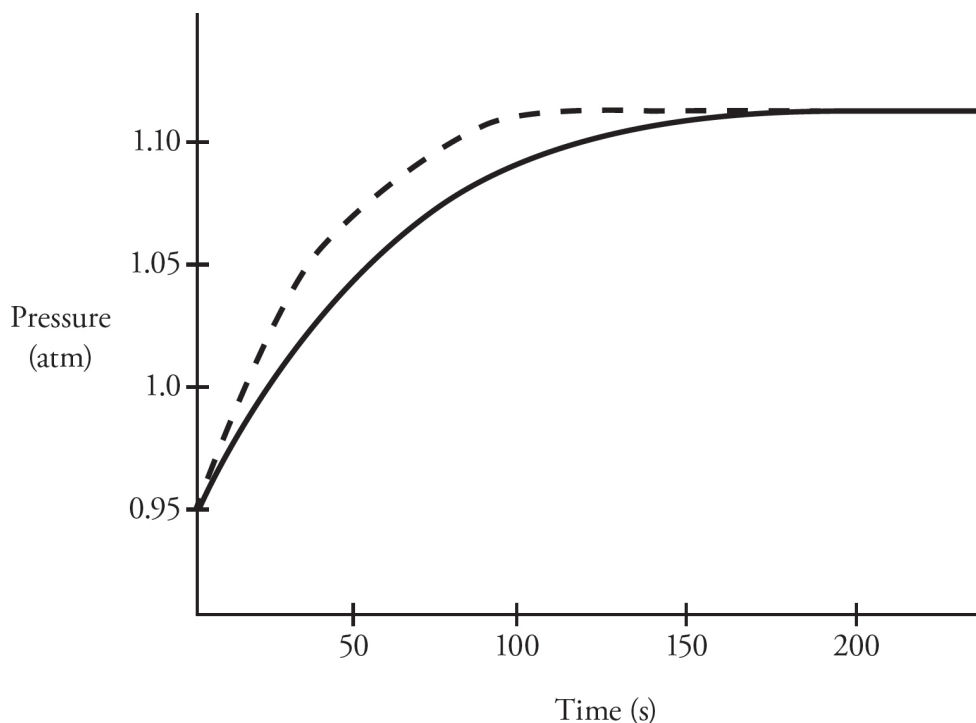
- (d) Catalysts are often used to lower the activation energy for a given reaction. The experiment is repeated, this time with a MnO_2 catalyst added.

- (i) How do catalysts lower a reaction's activation energy?

Catalysts lower activation energy by providing an alternative reaction pathway (a different set of elementary steps by which the reaction occurs).

- (ii) Using a dashed line, sketch a line on the graph that would indicate any changes that would occur in the data gathered during this trial.

The catalyst would not change the value for K_p , so the pressure would be the same. However, equilibrium would be established faster.



(e) Calculate the value for ΔG for this reaction at 23.0°C .

$$\Delta G = -RT \ln K \quad \Delta G = (-8.31 \text{ J/mol K})(296 \text{ K})(\ln 4.1 \times 10^{-3}) \quad \Delta G = +14,000 \text{ J}$$

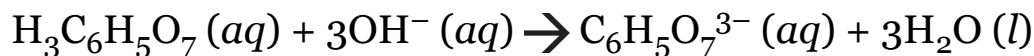
(f) Is the entropy change that occurs during this reaction positive or negative? Justify your answer.

The entropy change is positive. The reaction begins with a solid, and ends with a solid and a gas. The gas is much more disordered than the solid, leading to an increase in entropy.

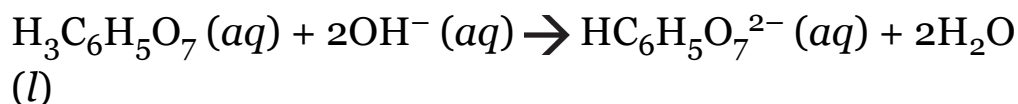
4. The primary acid in lemonade is citric acid ($\text{H}_3\text{C}_6\text{H}_5\text{O}_7$, $\text{p}K_a = 3.08$). To control the pH of many commercially available lemonade drinks, some sodium dihydrogen citrate ($\text{NaH}_2\text{C}_6\text{H}_5\text{O}_7$) is added to create a buffer system.

- (a) Write out the reaction which would occur if a strong base were added to a buffered lemonade solution such as the one described above.

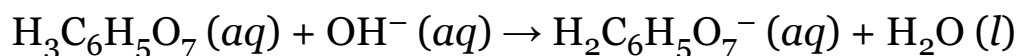
The exact reaction depends on how much strong base is added. Any of the following three answers would be acceptable.



OR



OR



- (b) What is the mol ratio of $\text{H}_2\text{C}_6\text{H}_5\text{O}_7^-$: $\text{H}_3\text{C}_6\text{H}_5\text{O}_7$ in a buffered solution with a pH of 3.25?

Using Henderson Haselbalch:

$$\text{pH} = \text{p}K_{\text{a}} + \log \frac{[\text{C}_6\text{H}_5\text{O}_7^{3-}]}{[\text{H}_3\text{C}_6\text{H}_5\text{O}_7]}$$

$$3.25 = 3.08 + \log \frac{[\text{C}_6\text{H}_5\text{O}_7^{3-}]}{[\text{H}_3\text{C}_6\text{H}_5\text{O}_7]}$$

$$0.17 = \log \frac{[\text{C}_6\text{H}_5\text{O}_7^{3-}]}{[\text{H}_3\text{C}_6\text{H}_5\text{O}_7]}$$

$$0.67 = \frac{[\text{C}_6\text{H}_5\text{O}_7^{3-}]}{[\text{H}_3\text{C}_6\text{H}_5\text{O}_7]}$$

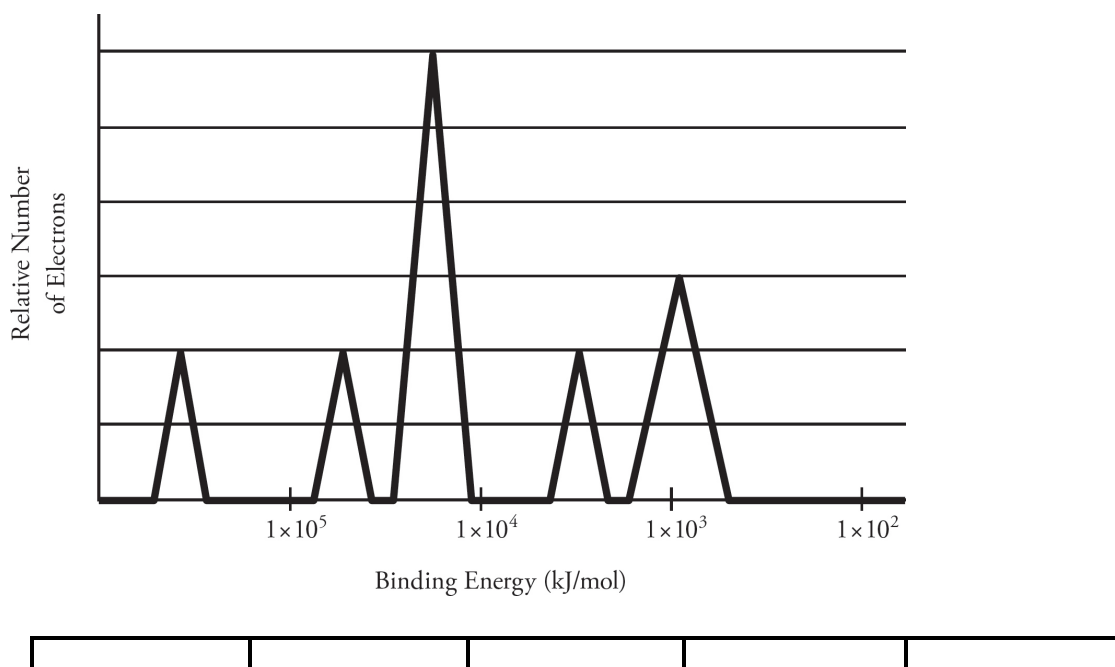
The mole ratio would be the same as the concentration ratio calculated above, as the volume of the solution is constant.

- (c) If a buffered solution of lemonade were to be diluted, would that increase the pH, decrease it, or leave it the same? Justify your answer.

There's two ways to answer this; and either would be acceptable as long as it is justified correctly. The textbook answer is that the pH would not change. This is because the ratio of the conjugate base:acid does not change, and in Henderson-Haselbalch if that ratio doesn't change, the pH of the buffer does not change.

That being said, if the buffer is diluted enough, the auto-dissociation of water may become a larger contributor to the solution acidity, and since water has a pH of 7 at 25°C adding more water to a buffered solution may cause the pH to rise slightly as a result.

5.



Peak 1	Peak 2	Peak 3	Peak 4	Peak 5
2.7×10^5 kJ/mol	5.4×10^4 kJ/mol	2.1×10^4 kJ/mol	4.9×10^3 kJ/mol	1.0×10^3 kJ/mol

The photoelectron spectrum for a neutral atom is displayed above.

- (a) What is the total number of valence electrons shown on the above spectrum?

The peaks, from left to right, are $1s$, $2s$, $2p$, $3s$, and $3p$. Valence electrons are those on the outmost principal energy level, which in this case is the third energy level. Looking at the $3s$ and $3p$ peaks, they represent a total of five electrons.

- (b) How much energy does a single photon of the radiation have?

First, convert the ionization energy from the outermost electron from a mole of electrons to a single electron:

$$\frac{1000 \text{ kJ}}{1 \text{ mol}} \times \frac{1000 \text{ J}}{1 \text{ kJ}} \times \frac{1 \text{ mol}}{6.02 \times 10^{23} \text{ photons}} = 1.66 \times 10^{-18} \text{ J}$$

For photoelectron spectroscopy, the incoming energy goes into two places. First, it overcomes the binding energy of the electron. Any energy left over after that goes into the kinetic energy of the electron once it is ejected. So:

$$\text{Incoming Energy} = \text{Binding Energy} + \text{Kinetic Energy}$$

$$4.17 \times 10^{-18} \text{ J} = 1.66 \times 10^{-18} \text{ J} + \text{KE}$$

$$\text{KE} = 2.51 \times 10^{-18} \text{ J}$$

(c) How much kinetic energy would the ejected electron have?

First, convert the energy from a single photon into kJ/mol.

$$\frac{4.17 \times 10^{-18} \text{ J}}{\text{photon}} \times \frac{6.02 \times 10^{23} \text{ photons}}{1 \text{ mol photons}} \times \frac{1 \text{ kJ}}{1000 \text{ J}} = 2510 \text{ kJ/mol}$$

For photoelectron spectroscopy, the incoming energy goes into two places. First, it overcomes the binding energy of the electron. Any energy left over after that goes into the kinetic energy of the electron once it is ejected. We can see from the data table that the valence (3p) electrons from phosphorus have a binding energy of 1000 kJ/mol. So:

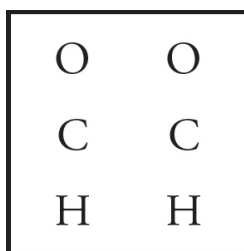
$$\text{Incoming Energy} = \text{Binding Energy} + \text{Kinetic Energy}$$

$$2510 \text{ kJ/mol} = 1000 \text{ kJ/mol} + \text{KE}$$

$$\text{KE} = 1510 \text{ kJ/mol}$$

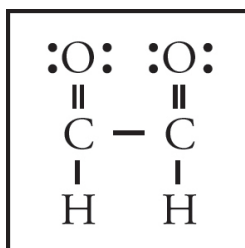
6. Glyoxal is an organic compound commonly used in many synthesis reactions.

(a) In the box below, complete the Lewis electron-dot diagram for the glyoxal molecule by drawing in all electron pairs



Glyoxal has a total of 22 valence electrons to place. If all single bonds are used, 26 electrons are placed. So, two double bonds are needed. Hydrogen cannot form double bonds, so that's out. If you double bond the two carbons

and one carbon-oxygen pair, that would give the carbon ten electrons, which is impossible because carbon cannot expand its octet. Thus, the only workable diagram has both carbon-oxygen pairs double bonded.



- (b) Identify all types of intermolecular forces that are present in a solution of glyoxal.

London dispersion forces are present, as are dipole-dipole attractions. Note that there are no hydrogen bonds, as for that to occur the hydrogen would need to be directly bonded to the oxygen, which it is not.

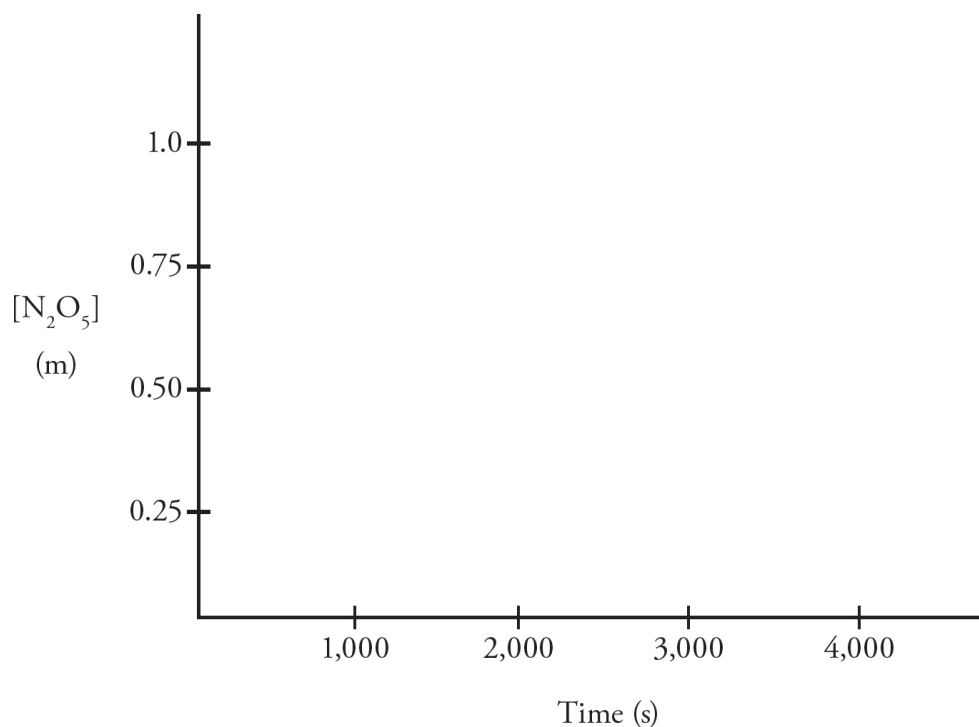
- (c) Would you expect glyoxal to be miscible with water? Justify your answer.

Miscibility occurs when two liquids have similar intermolecular forces. Water is very polar due to hydrogen bonding, and even though glyoxal does not have hydrogen bonding it is still polar, meaning it should be miscible in water.

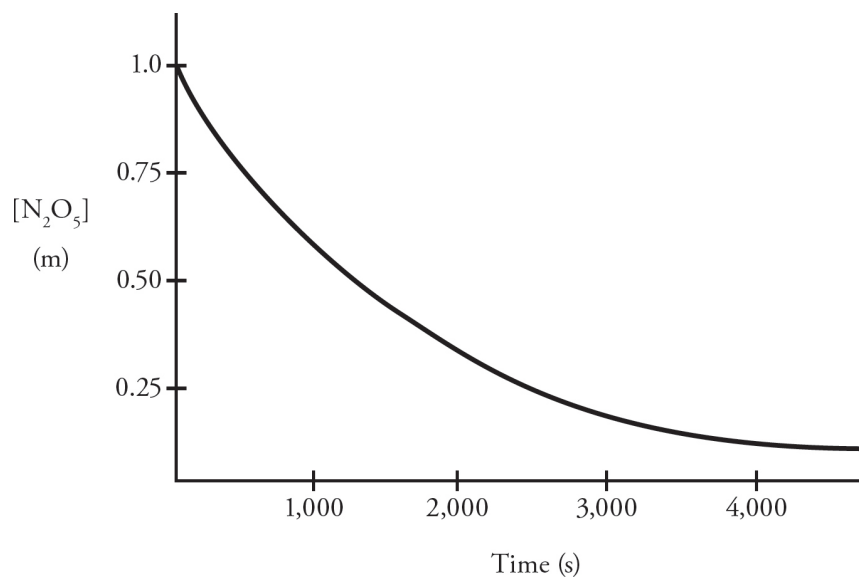


7. The decomposition of N_2O_5 is a first order reaction that goes to completion. A sample of N_2O_5 gas with a concentration of 1.0 M is placed in a sealed container at 50°C and allowed to decompose. The half-life of the reaction at 50°C is 1200 seconds.

- (a) On the graph below, draw a graph showing how the concentration of the N_2O_5 sample will change over time.



A first order reaction has a constant half life. So, ever 1200 s, half of the sample will have decayed. At 1200 s, the concentration would be 0.50 M. At 2400 s, the concentration would be 0.25 M, and so on.



(b) Calculate the rate constant for this reaction at 50°C

$$t_{\frac{1}{2}} = \frac{0.693}{k} \quad 1200 = \frac{0.693}{k} \quad k = 5.8 \times 10^{-4}$$

Note that the rate constant still needs units. The rate law is $\text{rate} = k[\text{N}_2\text{O}_5]$. Rate is measured in M/s, and concentration is measured in M. Plugging those units into the rate law yields: $\text{M/s} = k(\text{M})$ $k = \text{s}^{-1}$

So, the final answer is $5.8 \times 10^{-4} \text{ s}^{-1}$

(c) Using your graph, determine the instantaneous rate of disappearance for N_2O_5 at $t = 2000 \text{ s}$.

The integrated rate law for a first order reaction is $\ln[\text{N}_2\text{O}_5]_t = -kt + \ln[\text{N}_2\text{O}_5]_0$. Plugging in what we know:

$$\ln[\text{N}_2\text{O}_5]_{2000} = -(5.8 \times 10^{-4} \text{ s}^{-1})(2000 \text{ s}) + \ln(1.0 \text{ M})$$

$$\ln[\text{N}_2\text{O}_5]_{2000} = 1.08 \text{ M}$$

$$[\text{N}_2\text{O}_5]_{2000} = 0.34 \text{ M}$$

Once the concentration at $t = 2000 \text{ s}$ is known, then it can be plugged into the rate law to determine the rate at that point.

$$\text{Rate} = k[\text{N}_2\text{O}_5]$$

$$\text{Rate} = (5.8 \times 10^{-4} \text{ s}^{-1})(0.34 \text{ M})$$

$$\text{Rate} = 1.9 \times 10^{-4} \text{ M/s}$$

A stylized, abstract background featuring a network of overlapping gray lines that form a globe-like structure. The lines are of varying thickness and intersect to create a sense of depth and global connectivity.

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